

# Influence of dynamic ozone dry deposition on ozone pollution

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## Key Points:

- Remote and local ozone depositional sinks shape regional winter ozone pollution
- Dynamic ozone dry deposition changes summer surface ozone over northern mid-latitude regions by -4 to +7 ppb
- Variability and 21<sup>st</sup>-century changes in both stomatal and nonstomatal deposition influence summer surface ozone distributions

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**Abstract**

Identifying the contributions of chemistry and transport to observed ozone pollution using regional-to-global models relies on accurate representation of ozone dry deposition. We use a recently developed configuration of the NOAA GFDL chemistry-climate model – in which the atmosphere and land are coupled through dry deposition – to investigate the influence of ozone dry deposition on ozone pollution over northern mid-latitudes. In our model, deposition pathways are tied to dynamic terrestrial processes, such as photosynthesis and water cycling through the canopy and soil. Small increases in winter deposition due to more process-based representation of snow and deposition to surfaces reduce hemispheric-scale ozone through the lower troposphere by 5-12 ppb, improving agreement with observations relative to a simulation with the standard configuration for ozone dry deposition. Declining snow cover by the end of the 21<sup>st</sup> century tempers the previously identified influence of rising methane on winter ozone. Dynamic dry deposition changes summer surface ozone by -4 to +7 ppb. While previous studies emphasize the importance of uptake by plant stomata, new diagnostic tracking of depositional pathways reveals a widespread impact of nonstomatal deposition on ozone pollution. Daily variability in both stomatal and nonstomatal deposition contribute to daily variability in ozone pollution. 21<sup>st</sup>-century changes in summer deposition result from a balance among changes in individual pathways, reflecting differing responses to both high carbon dioxide (through plant physiology versus biomass accumulation) and water availability. Our findings highlight a need for constraints on the processes driving ozone dry deposition to test representation in regional-to-global models.

**1 Introduction**

In the troposphere, ozone is an air pollutant, a potent greenhouse gas, and an important source of the hydroxyl radical, the main tropospheric oxidant. Regional-to-global atmospheric chemistry models are key tools for quantifying the impacts of ozone pollution on human and vegetation health and pinpointing the drivers of observed trends and variability in tropospheric constituents. Representing ozone sources and sinks accurately in these models is fundamental to their utility. Ozone dry deposition is an important (20% of the annual global tropospheric loss), but uncertain and frequently overlooked, tropospheric ozone sink (Wild, 2007; Hardacre et al., 2015). Here we investigate the role of ozone dry deposition on ozone pollution at northern mid-latitudes with a global chemistry-climate model that leverages the carbon and water cycling in its underlying dynamic vegetation land model for representing dry deposition.

Dry deposition of ozone occurs through surface-mediated reactions after diffusion through plant stomata, or on leaf cuticles, other plant material, soil, water and snow. Ozone deposition velocity (a measure of the efficiency of the removal independent from ambient ozone concentration) is typically highest during summer, reflecting uptake by vegetation. Winter ozone dry deposition is usually not a research focus due to relatively low ozone deposition velocity. However, the long winter ozone lifetime implies efficient transport through large-scale circulation patterns, such that ozone at any particular location depends on both local and remote sources and sinks and thus may be sensitive to changes in ozone dry deposition locally and upwind. Although previous studies examine the sensitivity of winter ozone to ozone deposition velocity over the Uintah basin in the western United States (Matichuk et al., 2017) and boreal and Arctic regions (Helmig et al., 2007), it is unknown how ozone dry deposition impacts large-scale winter ozone over northern mid-latitudes. While ozone pollution is typically regarded as a summer problem (at least over polluted and populated regions), projected changes in anthropogenic precursor emissions drive large 21<sup>st</sup>-century increases in winter ozone (Clifton et al., 2014; Gao et al., 2013; Rieder et al., 2018), implying a need to advance understanding of winter ozone sources and sinks.

89 Much of the attention around ozone dry deposition is on its influence on summer  
90 ozone pollution. Previous work examines changes in ozone dry deposition with environ-  
91 mental conditions, ambient carbon dioxide, and land use/land cover as well as the im-  
92 pact of dry deposition on summer surface ozone (Solberg et al., 2008; Andersson & En-  
93 gardt, 2010; Ganzeveld et al., 2010; S. Wu et al., 2012; Trail et al., 2015; Fu & Tai, 2015;  
94 Huang et al., 2016; Geddes et al., 2016; Hollaway et al., 2016; Heald & Geddes, 2016;  
95 Anav et al., 2018; M. Lin et al., 2019; Wong et al., 2019). The aforementioned analy-  
96 ses linking surface ozone with ozone dry deposition all rely on models. These models typ-  
97 ically assume that stomatal uptake dominates ozone dry deposition and that nonstom-  
98 atal deposition is roughly constant or simply varies with leaf area index. However, lab-  
99 oratory and field evidence suggests that these assumptions may limit our ability to model  
100 ozone dry deposition accurately (Fuentes et al., 1992; Massman, 2004; Altimir et al., 2006;  
101 Cieslik, 2009; Fowler et al., 2009; Fares et al., 2010, 2012, 2014; Rannik et al., 2012; Potier  
102 et al., 2015, 2017; Sun et al., 2016; Clifton et al., 2017; Fumagalli et al., 2016; Clifton  
103 et al., 2019; Stella, Loubet, et al., 2011). Current understanding of nonstomatal depo-  
104 sition pathways is that leaf cuticular uptake increases with leaf wetness, soil uptake de-  
105 creases with soil moisture, and snow on vegetation and the ground decreases uptake (Clifton  
106 et al., 2020). Systematic omissions in process representation that lead to variations in  
107 ozone deposition velocity with meteorology or biophysics may impede accurate model  
108 simulations of changes in ozone pollution attributable to changes in dry deposition.

109 Here we probe the influence of ozone dry deposition on winter and summer ozone  
110 pollution over northern mid-latitudes under a 21<sup>st</sup>-century scenario for climate and an-  
111 thropogenic precursor emissions using a new configuration of the global National Oceanic  
112 and Atmospheric Administration (NOAA) Geophysical Fluid Dynamics Laboratory (GFDL)  
113 chemistry-climate model. In particular, we use the biophysics of the land component to  
114 simulate ozone dry deposition by plant stomata, stems, and wet, dry, and snow-covered  
115 soil and leaf cuticles. We evaluate this model with ozone eddy covariance flux observa-  
116 tions from long-term and short-term datasets and estimates of the stomatal fraction of  
117 ozone dry deposition derived from observations. We compare simulations with this new  
118 dynamic ozone dry deposition scheme to simulations using a prescribed climatology of  
119 ozone deposition velocity, the default configuration in the GFDL model. While nonstom-  
120 atal deposition pathways represent observed dependencies on meteorological and biophys-  
121 ical variables in our model to the extent possible, these pathways remain uncertain due  
122 to a paucity of observational constraints, and their representation in models is highly pa-  
123 rameterized (Clifton et al., 2020). Our goal is to investigate how dynamic ozone dry de-  
124 position, based on current understanding, influences ozone pollution.

## 125 2 Methods

126 We conduct time-slice simulations for the 2010s and 2090s with the NOAA GFDL  
127 atmospheric model version 3 (AM3) coupled to the NOAA GFDL land model version  
128 3 (LM3) through not only carbon, water, and energy exchanges but also dry deposition  
129 of several atmospheric constituents (AM3DD) (Paulot et al., 2018). Each simulation con-  
130 tains ten years. Below we describe the model configuration and the dynamic dry depo-  
131 sition scheme for ozone, which we modify from the general dynamic dry deposition scheme  
132 described by Paulot et al. (2018).

133 AM3 is a chemistry-climate model with online fully coupled stratospheric and tropo-  
134 spheric chemistry (Naik et al., 2013; Donner et al., 2011). We use AM3 with C48 (cubed  
135 sphere) configuration (approximately 2° by 2°) and 48 vertical levels. We update the treat-  
136 ment of wet deposition of aerosols and gases in AM3 following Paulot et al. (2016); in  
137 particular, snow formed by the Bergeron process does not scavenge water-soluble aerosols.

138 We use Representative Concentration Pathway 8.5 (RCP8.5) (van Vuuren et al.,  
139 2011; Riahi et al., 2011; Lamarque et al., 2011), the high-warming scenario designed for

140 the Coupled Model Intercomparison Project 5, to represent 21<sup>st</sup>-century climate and an-  
 141 thropogenic emissions. Aerosol and ozone precursor emissions and global concentrations  
 142 of greenhouse gases are set to 2010 and 2090 levels for our 2010s and 2090s time-slice  
 143 simulations, respectively. Isoprene emissions are calculated online with a version of Model  
 144 of Emissions of Gases and Aerosols from Nature (MEGAN) in AM3 (Guenther et al.,  
 145 2006; Emmons et al., 2010; Rasmussen et al., 2012). Simulations are forced with decadal  
 146 mean (2011-2020 or 2091-2100) sea ice and sea-surface temperatures from transient RCP8.5  
 147 simulations (average over three ensemble members) from the NOAA GFDL coupled model  
 148 version 3. We use initial conditions for 2010 and 2090 from one ensemble member of the  
 149 transient 21<sup>st</sup>-century RCP8.5 simulations described in Clifton et al. (2014) that were  
 150 spun up from a pre-industrial control simulation (John et al., 2012).

151 LM3 is a global land model with terrestrial carbon, energy and water cycling, dy-  
 152 namic vegetation, and land use transitions (Shevliakova et al., 2009; Milly et al., 2014).  
 153 A sub-grid tiling framework in LM3 allows individual tiles to represent distinct land uses,  
 154 including primary vegetation, cropland, pasture, secondary vegetation, as well as bod-  
 155 ies of water and glaciers. We prescribe land use distributions with either 2010 or 2090  
 156 RCP8.5 (Hurtt et al., 2011). Primary vegetation has never been disturbed by humans  
 157 directly, whereas secondary vegetation has been harvested and subsequently abandoned  
 158 at least once. Each grid cell contains up to twelve stages of secondary vegetation, allow-  
 159 ing for differing recovery times. Modifications to crop harvesting and pasture grazing fol-  
 160 low Paulot et al. (2018). Each vegetated sub-grid tile has one land cover type. Land cover  
 161 types include temperate deciduous forests, tropical forests, coniferous forests, C<sub>3</sub> grass,  
 162 and C<sub>4</sub> grass. The distribution of vegetation evolves with climate, but the distribution  
 163 of bodies of water and glaciers is time invariant. There are five pools of vegetation biomass  
 164 (leaves, fine roots, sapwood, heartwood, and labile stores), and allocation rules and daily  
 165 net primary production update the pools each day (Shevliakova et al., 2009). Phenol-  
 166 ogy (*i.e.*, leaf on/off) and thus leaf area index (LAI) is updated monthly from the leaf  
 167 biomass pool according to monthly mean air temperature and soil water available to the  
 168 plant (Shevliakova et al., 2009) except for temperate deciduous vegetation, for which LAI  
 169 has strong seasonality. We update the temperate deciduous vegetation daily according  
 170 to critical temperature and growing degree day following Weng et al. (2015).

## 171 2.1 Ozone dry deposition in AM3DD

172 The new ozone dry deposition parameterization in LM3 uses a big-leaf resistance  
 173 framework. Pathways for ozone dry deposition include leaf cuticles, stomata, stems, and  
 174 the ground. Ozone deposition velocity ( $v_d$ ) [cm s<sup>-1</sup>] follows:

$$175 \quad v_d = \left[ R_a + \frac{1}{\frac{1}{R_{b,v} + \frac{1}{\frac{1}{R_{cut}} + \frac{1}{R_{stom} + R_{meso}}}} + \frac{1}{R_{b,v} + R_{stem}} + \frac{1}{R_{ac,g} + R_{b,g} + R_g}} \right]^{-1} * 100 \quad (1)$$

176 In the following paragraphs, we define each resistance term in equation 1. The scheme  
 177 follows Paulot et al. (2018) except where otherwise noted.

178 The resistance to turbulent transport between the atmosphere and canopy ( $R_a$ )  
 179 [s m<sup>-1</sup>] follows Fick's Law and Monin-Obukhov Similarity Theory. The quasi-laminar  
 180 boundary-layer resistance for vegetation ( $R_{b,v}$ ) [s m<sup>-1</sup>] follows Choudhury and Monteith  
 181 (1988):

$$182 \quad R_{b,v} = \frac{a}{b} \sqrt{\frac{d_{leaf}}{u_h}} \left[ 1 - e^{-a/2} \right] \left( \frac{Sc}{Pr} \right)^{2/3} \quad (2)$$

183  $d_{leaf}$  is the leaf dimension [m];  $u_h$  is wind speed at the top of the canopy [m s<sup>-1</sup>] ( $h$  is  
 184 canopy height [m]);  $a$  is an empirical constant (value of 3);  $b$  [m s<sup>-0.5</sup>] is an empirical  
 185 constant (value of 0.02);  $Sc$  is the Schmidt number [unitless];  $Pr$  is the Prandtl number



186 [unitless].  $R_{b,v}$  is scaled by fraction of vegetation that is stems versus leaves when used  
 187 in equation 1.

188 Paulot et al. (2018) apply both equation 2 and the Jensen and Hummelshøj (1995,  
 189 1997)  $R_{b,v}$  parameterization, with the intention of including a resistance to in-canopy  
 190 turbulence. However, equation 2 is a quasi-laminar boundary-layer resistance, not a re-  
 191 sistance to in-canopy turbulence. We use equation 2 for  $R_{b,v}$  because it is used for en-  
 192 ergy and carbon exchanges in LM3. A resistance to in-canopy turbulence for leaf depo-  
 193 sition is unnecessary in our big-leaf model because  $R_a$  accounts for turbulent transport  
 194 between the atmosphere and canopy and all vegetation is assumed to be at the canopy  
 195 height.

196 We distinguish cuticular deposition among dry, wet, and snow-covered leaves. Frac-  
 197 tional leaf wetness is calculated from canopy-intercepted water, specifically the ratio of  
 198 canopy-intercepted water to the maximum storage capacity to the two-thirds power (Bonan,  
 199 1996). Fractional snow cover on vegetation is calculated in the same way but with canopy-  
 200 intercepted snow. We employ an adjustment function  $s$  [unitless] to reduce wet and dry  
 201 cuticular deposition when leaf temperatures are cold ( $<5^\circ\text{C}$ ).

$$202 \quad s(T_{leaf}) = \max[e^{-c(T_{leaf}-5)}, 1] \quad (3)$$

203  $T_{leaf}$  is leaf temperature [ $^\circ\text{C}$ ];  $c$  is a constant [ $^\circ\text{C}^{-1}$ ]. Such an adjustment function as-  
 204 sumes that the chemistry on surfaces is slower when the surfaces are cold. We use  $c=0.9$   
 205  $^\circ\text{C}^{-1}$  for wet and  $c=0.1$   $^\circ\text{C}^{-1}$  for dry cuticular deposition, employing different values  
 206 because the initial resistances for wet and dry cuticular deposition differ by an order of  
 207 magnitude (see below). Our temperature adjustment function, an adaptation of Zhang  
 208 et al. (2003), allows for cuticular deposition at cold temperatures to be reduced, but not  
 209 turned off. We do not turn off cuticular deposition on cold surfaces following observa-  
 210 tional evidence that uptake occurs on material protruding from snow (Clifton et al., 2020).

211 Paulot et al. (2018) use the Zhang et al. (2003) temperature adjustment function.  
 212 Without our change to the Zhang et al. (2003) temperature adjustment function, win-  
 213 ter cuticular uptake to coniferous forests (only in boreal regions in LM3) becomes higher  
 214 than supported by field observations. For example, simulated winter mean  $v_d$  over bo-  
 215 real regions ( $55\text{-}65^\circ\text{N}$ ) with  $\text{LAI} \geq 2 \text{ m}^2 \text{ m}^{-2}$  is  $0.1 \text{ cm}^{-1}$  with this temperature adjust-  
 216 ment function, only slightly less than observations from Hyytiälä, a boreal coniferous for-  
 217 est, which suggest a winter mean  $v_d$  of  $0.12 \text{ cm s}^{-1}$ . Previous studies do not identify the  
 218 need for a stronger temperature adjustment function, likely because they assume win-  
 219 ter boreal regions are completely snow-covered, whereas here we consider dynamic canopy  
 220 cycling of snow. Canopy snow cycling in LM3 allows conifers to be occasionally snow-  
 221 free, leading us to implement a stronger temperature adjustment function to reduce oth-  
 222 erwise unrealistically high simulated uptake to bare conifer cuticles.

223 The resistance to cuticular deposition to dry leaves ( $R_{cut,dry}$ ) [ $\text{s m}^{-1}$ ] follows:

$$224 \quad R_{cut,dry} = \frac{R_{i,cut,dry}}{\text{LAI}e^{\text{RH}}} s(T_{leaf}) \quad (4)$$

225  $R_{i,cut,dry}$  is the initial resistance to dry cuticular deposition [ $\text{s m}^{-1}$ ]; RH is fractional in-  
 226 canopy relative humidity [unitless]. The RH dependence is an update to Paulot et al.  
 227 (2018) and follows field and laboratory evidence suggesting that ozone dry deposition  
 228 to cuticles occurs through aqueous surface-mediated chemistry (Fuentes et al., 1992; Zhang  
 229 et al., 2002; Potier et al., 2015, 2017; Sun et al., 2016). In particular, the RH dependence  
 230 in the model for  $R_{cut,dry}$  represents the thin water films that form on leaves at high am-  
 231 bient humidity (Burkhardt & Hunsche, 2013).

232 Higher ozone deposition to leaves wet by rain and dew (Clifton et al., 2020) is also  
 233 accounted for in our model. The resistance to cuticular deposition to leaves wet by rain

234 and dew ( $R_{cut,wet}$ ) [ $s\ m^{-1}$ ] follows:

$$235 \quad R_{cut,wet} = \frac{R_{i,cut,wet}}{LAI} s(T_{leaf}) \quad (5)$$

236  $R_{i,cut,wet}$  is the initial resistance to wet cuticular deposition [ $s\ m^{-1}$ ]. For pastures, crops,  
 237 and grasses,  $R_{i,cut,dry}$  is  $4000\ s\ m^{-1}$  and  $R_{i,cut,wet}$  is  $200\ s\ m^{-1}$  and for coniferous, tem-  
 238 perate deciduous, and tropical trees,  $R_{i,cut,dry}$  is  $6000\ s\ m^{-1}$  and  $R_{i,cut,wet}$  is  $400\ s\ m^{-1}$ .  
 239 Initial resistances follow Zhang et al. (2003), except that initial resistances for conifer-  
 240 ous trees are the same for other trees, not much lower as suggested by Zhang et al. (2003).  
 241 Paulot et al. (2018) originally implemented the initial resistances suggested by Zhang  
 242 et al. (2003) for conifers, but increasing the initial resistances for conifers to agree with  
 243 the values for other trees reduces dry deposition to coniferous forests (only in boreal re-  
 244 gions in LM3) where LM3 overestimates LAI. We note that the Zhang et al. (2003) ini-  
 245 tial resistances were derived from observations from one growing season or less in east-  
 246 ern U.S. locations and thus their application more generally for global land use types is  
 247 highly uncertain.

248 The resistance to cuticular deposition to snow-covered leaves ( $R_{cut,snow}$ ) [ $s\ m^{-1}$ ]  
 249 follows:

$$250 \quad R_{cut,snow} = \frac{R_{i,snow}}{LAI} \quad (6)$$

251  $R_{i,snow}$ , the initial resistance to snow, is  $7000\ s\ m^{-1}$ . Often the number of surfaces cov-  
 252 ered by snow is not considered in models of ozone dry deposition (*i.e.*,  $R_{cut,snow} = R_{i,snow}$ ).  
 253 Our model (equation 6) assumes deposition increases with LAI [ $m^2\ m^{-2}$ ], implying more  
 254 deposition with a larger surface area covered with snow. This relationship is supported  
 255 by observations of relatively high  $v_d$  over snow-covered forests (Padro et al., 1992; Padro,  
 256 1993; Z. Wu et al., 2016; Neiryck & Verstraeten, 2018).

257 Our value for  $R_{i,snow}$  is more than triple the  $2000\ s\ m^{-1}$  used by Paulot et al. (2018)  
 258 and given by Zhang et al. (2003). Increasing  $R_{i,snow}$  leads to uptake by snow on the ground  
 259 and leaf cuticles of  $0.015\ cm\ s^{-1}$  on average over  $40-65^\circ N$  for present-day winter, agree-  
 260 ing with most field and laboratory observations supporting  $v_d$  for snow-covered regions  
 261 higher than  $0.01\ cm\ s^{-1}$  (Aldaz, 1969; Colbeck & Harrison, 1985; I. Galbally & Allison,  
 262 1972; I. E. Galbally & Roy, 1980; Wesely et al., 1981; Stocker et al., 1995; Gong et al.,  
 263 1997; Hopper et al., 1998; Helmig et al., 2009; Clifton et al., 2020).

264 Stomatal resistance ( $R_{stom}$ ) [ $s\ m^{-1}$ ] is calculated explicitly from net photosynthe-  
 265 sis ( $A_{net}$ ) [ $mol\ CO_2\ m^{-2}\ s^{-1}$ ] (Farquhar et al., 1980; Collatz et al., 1991, 1992) via Leuning  
 266 (1995):

$$267 \quad R_{stom} = \frac{p_s}{RT_{leaf}} \frac{1}{m} \left(1 + \frac{d_s}{d_0}\right) \frac{c_i - \Gamma}{A_{net}} LAI \quad (7)$$

268  $p_s$  is surface pressure [Pa];  $R$  is the universal gas constant [ $J\ mol\ air^{-1}\ K^{-1}$ ];  $m$  is an em-  
 269 pirical constant [unitless];  $d_s$  is the humidity deficit [ $kg\ H_2O\ kg\ air^{-1}$ ];  $d_0$  is another em-  
 270 pirical constant [ $kg\ H_2O\ kg\ air^{-1}$ ];  $c_i$  is carbon dioxide concentration internal to the leaf  
 271 [ $mol\ CO_2\ mol\ air^{-1}$ ];  $\Gamma$  is carbon dioxide compensation point [ $mol\ CO_2\ mol\ air^{-1}$ ];  $R_{stom}$   
 272 shown in the above equation is also scaled by the inverse of the fractional water stress  
 273 if the fractional water stress  $< 1$  (Milly et al., 2014). The water stress is the ratio of wa-  
 274 ter supply to roots to water demand from atmosphere.

275 We account for the different diffusivities of ozone and water vapor by scaling  $R_{stom}$   
 276 given in equation 7 for water vapor by the ratio of the diffusivities of the two gases. The  
 277 resistance to ozone reacting with internal fluids and tissues in our model (*i.e.*, often called  
 278 a mesophyll resistance, or  $R_{meso}$  [ $s\ m^{-1}$ ]) is small ( $0.01\ s\ m^{-1}$ ) because laboratory ev-  
 279 idence suggests that ozone reacts immediately upon entering stomata (Laisk et al., 1989;  
 280 D. Wang et al., 1995).

281 Stomatal deposition is reduced on the part of the leaf that is wet by dew or rain;  
 282 this happens through a 30% decrease in  $A_{net}$  and stomatal conductance on the wet part

283 of the leaf. This is a correction to Paulot et al. (2018) and M. Lin et al. (2019) who  
 284 reduce stomatal deposition by the fraction of the leaf that is wet in addition to the 30%  
 285 decrease in  $A_{net}$  and stomatal conductance that we retain here.

286 The stem resistance ( $R_{stem}$ ) to ozone dry deposition is:

$$287 \quad R_{stem} = \frac{R_{i,stem}}{SAI} \quad (8)$$

288  $R_{i,stem}$  is  $3000 \text{ s m}^{-1}$ ; SAI [ $\text{m}^2 \text{ m}^{-2}$ ] is stem area index. While Paulot et al. (2018) use  
 289  $4000 \text{ s m}^{-1}$  and the Zhang et al. (2003) temperature adjustment function to reduce stem  
 290 deposition onto cold surfaces, our change to  $R_{i,stem}$  and removal of the temperature ad-  
 291 justment function allow for higher deposition to stems and distinguishing between winter  
 292 deposition to vegetated versus non-vegetated regions (Clifton et al., 2020). The lat-  
 293 ter also allows for slightly higher winter deposition to bare deciduous trees relative to  
 294 areas without woody biomass, as supported by observations (Padro et al., 1992; Clifton  
 295 et al., 2020).

296 A resistance to in-canopy turbulence influences dry deposition to the ground when  
 297 vegetation is present ( $LAI+SAI > 0.25 \text{ m}^2 \text{ m}^{-2}$ ) and follows Paulot et al. (2018). The  
 298 model was developed from a very short-term regression analysis over a corn field (Van  
 299 Pul & Jacobs, 1994), but has been used widely in dry deposition schemes (Erisman et  
 300 al., 1994; Zhang et al., 2002; Emberson et al., 2000; Pleim & Ran, 2011). We use this  
 301  $R_{ac,g}$  parameterization because there are not many alternatives for large-scale big-leaf  
 302 modeling.

$$303 \quad R_{ac,g} = \frac{14(LAI + SAI)h}{u_*} \quad (9)$$

304  $u_*$  is friction velocity [ $\text{m s}^{-1}$ ]. The number 14 is a constant fit via regression and has units  
 305 of  $\text{m}^{-1}$ . Instead of setting LAI to unity when trees are leafless as Erisman et al. (1994)  
 306 do, we replace LAI with LAI+SAI for all conditions. If vegetation is not present,  $R_{ac,g}$   
 307 is negligible ( $0.01 \text{ s m}^{-1}$ ).

308 The quasi-laminar boundary-layer resistance for all ground surfaces ( $R_{b,g}$ ) [ $\text{s m}^{-1}$ ]  
 309 except lakes follows Wesely and Hicks (1977) implemented by Paulot et al. (2018):

$$310 \quad R_{b,g} = \frac{2}{k u_*} \left( \frac{Sc}{Pr} \right)^{2/3} \quad (10)$$

311  $k$  is the von Kármán constant [unitless]. If vegetation is present then  $u_*$  near the ground  
 312 ( $u_{*,g}$ ) [ $\text{m s}^{-1}$ ] is used in equation 10.

$$313 \quad u_{*,g} = u_* e^{0.6(LAI+SAI)\left(\frac{z_{0,g}}{h}-1\right)} \quad (11)$$

314  $z_{0,g}$  is the roughness length of the ground for scalars [m] as calculated in Bonan (1996).  
 315 Equation 11 follows Loubet et al. (2006) but also includes SAI, allowing bare trees to  
 316 contribute to drag. For very low vegetation ( $h < 0.1 \text{ m}$ ), we assume  $u_{*,g} = u_*$ .

317 The quasi-laminar boundary-layer resistance for lakes ( $R_{b,g,lake}$ ) [ $\text{s m}^{-1}$ ] follows Hicks  
 318 and Liss (1976):

$$319 \quad R_{b,g,lake} = \frac{\ln\left(\frac{z_{0,g}}{D_{O_3} k u_*}\right)}{k u_*} \quad (12)$$

320  $D_{O_3}$  is the diffusivity of ozone in air [ $\text{m}^2 \text{ s}^{-1}$ ].

321 We distinguish dry deposition to the ground among snow-covered, wet, and dry soil,  
 322 deserts, lakes, and glaciers. While a synthesis across observations suggests ground de-  
 323 position depends on soil moisture (Massman, 2004), the exact relationship is unknown.  
 324 We thus prescribe a simple step function such that ground uptake decreases when soil

is wet as suggested by Massman (2004). We define wet soil as fractional surface soil moisture in a tile  $>0.9$ . Some work points to an exponential or logarithmic dependence of ground deposition with moisture (Stella, Loubet, et al., 2011; Stella et al., 2019; Fumagalli et al., 2016), but we maintain a simpler change in ground deposition due to poor understanding of what happens at the large scale.

The treatment of ground deposition to cold surfaces from Paulot et al. (2018) considers the ground to be covered with snow if there is any snow in a tile and employs the Zhang et al. (2003) temperature adjustment function to reduce ground deposition at cold temperatures. Instead, we update the model to use fractional snow cover on the ground, calculated as a function of snow depth and prescribed critical depth as done for surface albedo. We change the temperature adjustment function to the one used for cuticles (equation 3) and use  $c=0.025 \text{ } ^\circ \text{C}^{-1}$  and soil temperature ( $T_{soil}$ ) [ $^\circ \text{C}$ ]. We maintain the Paulot et al. (2018) treatment of frozen lakes: lakes are frozen if there is any solid water.

The resistance to ground deposition ( $R_g$ ) [ $\text{s m}^{-1}$ ] follows:

$$R_g = R_{i,g} s(T_{soil}) \quad (13)$$

$R_{i,g}$  [ $\text{s m}^{-1}$ ] is the initial resistance to ground deposition.  $R_{i,g}$  for snow and ice is  $R_{i,snow}$  ( $7000 \text{ s m}^{-1}$ ).  $R_{i,g}$  for wet surfaces (*e.g.*, lakes, wet soil) is  $500 \text{ s m}^{-1}$  and dry vegetated surfaces is  $200 \text{ s m}^{-1}$  (Zhang et al., 2003).  $R_{i,g}$  for deserts (defined by  $<0.05 \text{ kg m}^{-2}$  biomass) is  $500 \text{ s m}^{-1}$ . Ozone dry deposition to the ground is largely considered to occur through reaction with soil organic material, but short-term observations suggest non-negligible uptake over the Sahara Desert (Güsten et al., 1996). However, relationships between soil organic content and ozone dry deposition to the ground are poorly constrained, leading to major uncertainties in representing dry deposition in different dry environments. Paulot et al. (2018) define  $R_{i,g}$  for deserts to be  $500 \text{ s m}^{-1}$ , but their desert definition is broader ( $<0.25 \text{ kg m}^{-2}$  biomass). Our changes to ground deposition to deserts in part reflect the need for non-negligible deposition in regions such as the western US where otherwise  $v_d$  in LM3 is too low due to inaccurate representation of vegetation there.

In order to probe the contribution of different deposition pathways to  $v_d$ , we examine effective conductances. Generally, a conductance is the inverse of a resistance. The effective conductance is the amount of deposition (in velocity units) occurring through a given deposition pathway. The sum of all of the effective conductances is  $v_d$ .

Dry deposition to the ocean in AM3DD follows monthly average fields from GEOS-Chem, a widely used chemical transport model. Aside from the meteorological dependencies of the resistances to turbulent transport and the quasi-laminar boundary layer between the ocean and atmosphere,  $v_d$  in GEOS-Chem over oceans does not change with meteorology, sea-surface temperatures, or surface-mediated chemistry in contrast to observational evidence (Ganzeveld et al., 2009; Martino et al., 2012; Helmig et al., 2012; Sarwar et al., 2016; Luhar et al., 2017).

## 2.2 Sensitivity simulation with default configuration for ozone dry deposition

In addition to AM3DD simulations with dynamic ozone dry deposition, we examine AM3DD simulations where we prescribe a monthly mean climatology of  $v_d$  scaled to a diel cycle (hereafter, AM3DD-staticO3DD), which is the default configuration for the GFDL model (Naik et al., 2013; Paulot et al., 2016). The climatology is single-year monthly average fields from a widely used chemical transport model, GEOS-Chem. We impose the multiyear monthly mean diel cycle from AM3DD 2010s so that differences between AM3DD and AM3DD-staticO3DD reflect differences in interannual, daily, and spatial variability and 21<sup>st</sup>-century changes in  $v_d$  rather than the diel cycle. AM3DD-staticO3DD for the 2090s uses the same setup for  $v_d$  as AM3DD-staticO3DD for the 2010s,

374 which allows us to consider how neglecting 21<sup>st</sup>-century changes in  $v_d$  impacts surface  
375 ozone projections.

376 Briefly, the  $v_d$  climatology was generated with GEOS-Chem, which uses a mod-  
377 ified Wesely (1989) dry deposition scheme (Y. Wang et al., 1998).  $R_a$  follows Fick’s Law  
378 and Monin-Obukhov Similarity Theory (specifically, Businger et al. (1971)) and  $R_b$  fol-  
379 lows Wesely and Hicks (1977).  $R_g$  and  $R_{ac,g}$  are time-invariant, but change with land  
380 cover type. Ozone dry deposition to cuticles varies with LAI and land cover type. Land  
381 cover type follows the Olson et al. (2001) land map. Stomatal ozone dry deposition varies  
382 with LAI, light, temperature, and land cover type (Y. Wang et al., 1998). This scheme  
383 also has a deposition pathway to the ground as well as to the lower canopy. High albedo  
384 ( $>0.4$ ) is used as a proxy for snow-covered surfaces to which ozone dry deposition is in-  
385 hibited. The temperature adjustment function for cold surfaces in GEOS-Chem follows  
386 Wesely (1989).

### 387 **3 Model evaluation of dynamic ozone dry deposition**

388 We compare monthly mean  $v_d$  from ozone eddy covariance fluxes at observational  
389 sites (Table 1) with  $v_d$  simulated by AM3DD (Figures 1, 2). We archive simulated  $v_d$   
390 for each land cover type within a grid cell (recall sub-tiling framework described above),  
391 which allows for a more direct comparison with observations (*e.g.*, Paulot et al. (2018),  
392 Silva and Heald (2018)). The model land cover type that best matches the observational  
393 site is selected for the evaluations in Figures 1 and 2. We focus our model evaluation on  
394 the eight sites with multiple years of data with at least a couple of months of data col-  
395 lected in a given year (Figure 1). At these sites, monthly daily mean  $v_d$  shows strong  
396 interannual variability, similar to that identified by Clifton et al. (2017) for monthly day-  
397 time mean  $v_d$  at Harvard Forest. For most sites, simulated  $v_d$  is close to the multiyear  
398 mean observed  $v_d$  and mostly within the observed range of interannual variability (Fig-  
399 ure 1). Two exceptions are the sites in Italy during nonsummer months – whereas AM3DD  
400 slightly overestimates  $v_d$  at Castelporziano, AM3DD slightly underestimates  $v_d$  at Is-  
401 pra. The model also slightly overestimates summer  $v_d$  at Grignon and winter  $v_d$  at Blod-  
402 gett Forest, suggesting that the model may struggle to capture  $v_d$  in Mediterranean-like  
403 ecosystems. Nonetheless, overall, we suggest that AM3DD captures observed  $v_d$  patterns  
404 on a climatological basis at long-term monitoring sites.

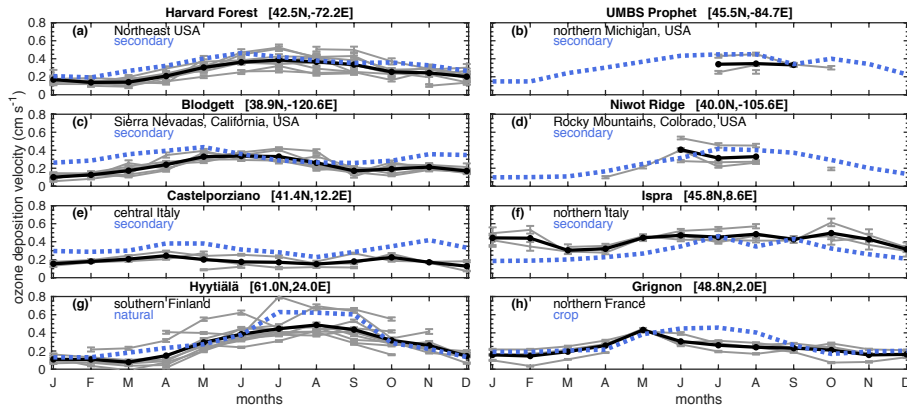
405 At the sites with shorter-term measurements, simulated monthly mean  $v_d$  tends  
406 to overestimate observed  $v_d$  (Figure 2a,b,c,d), except for Lincove, the orange orchard in  
407 the Central Valley of California, during nonspring months. In general, long-term ozone  
408 flux observations at these sites are necessary to understand the full extent of the appar-  
409 ent biases. We note that the short-term observations from Bondville, Kane, and Sand  
410 Flats were used in the development of the nonstomatal deposition parameterization from  
411 Zhang et al. (2002, 2003) from which we use some initial resistances. Agreement between  
412 simulated and observed  $v_d$  at these sites is lower relative to other sites, suggesting model  
413 performance does not follow implicit tuning.



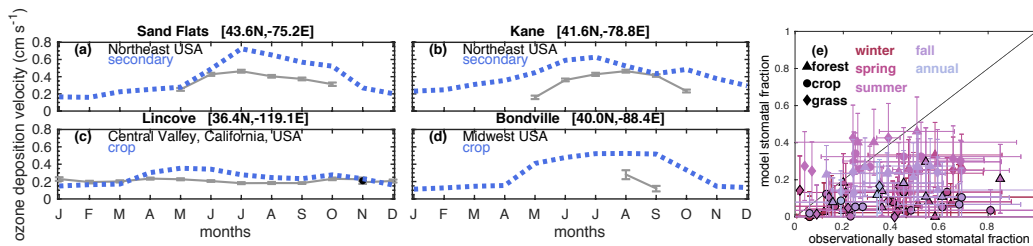
**Table 1.** Sites with ozone eddy covariance fluxes used in model evaluation

Site	Location	Land Cover	Year(s)	Previous References	Details
Blodgett Forest	38.9°N, 120.63°W	forest	2001-2006	Fares et al. (2010)	
Bondville	40.05°N, 88.37°W	crop	1994	Meyers et al. (1998); Finkelstein (2001)	1
Castelporziano	41.42°N, 12.21°E	forest	2012-2015	Fares et al. (2014)	
Grignon	48.84°N, 1.95°E	crop	2004-2008	Stella, Personne, et al. (2011)	
Harvard Forest	42.53°N, 72.18°W	forest	1990-2000	Munger et al. (1996); Clifton et al. (2017, 2019)	4,7,a
Hyytiälä	61.85°N, 24.28°E	forest	2002-2012	Altimir et al. (2006); Rannik et al. (2012)	2,8
Ispra	45.81°N, 8.63°E	forest	2013-2015		2,3,a
Kane Experimental Forest	41.59°N, 78.76°W	forest	1997	Meyers et al. (1998); Finkelstein et al. (2000)	1,4,5,7
Lincove Orange Orchard	36.36°N, 119.09°W	crop	2009-2010	Fares et al. (2012)	1,4
Niwot Ridge	40.03°N, 105.55°W	forest	2002-2005	Turnipseed et al. (2009)	
Sand Flats State Forest	43.565°N, 75.23°W	forest	1998	Meyers et al. (1998); Finkelstein et al. (2000)	1,4,7
UMBS Prophet	45.5°N, 84.7°W	forest	2002-2005	Hogg et al. (2007); Hogg (2007)	2,6,a

Different data filtering approaches were applied by the individual data providers; sometimes the datasets that we received were already filtered for outliers, sometimes not. Applying different filtering techniques for the datasets is our attempt to achieve an overall similar level of filtering among datasets. In the “Details” column, numbers indicate that we further filtered the data that we received: 1 is no  $|v_d| > 10 \text{ cm s}^{-1}$ ; 2 is no  $|v_d| > 5 \text{ cm s}^{-1}$ ; 3 is no level 2 values; 4 is no  $v_d$  outside  $\mu \pm 3\sigma$ ; 5 is we do not include missing half-hourly ozone fluxes at 23:30 local time many nights in July as missing data; 6 is no 2003 data after 5 September; 7 is  $v_d$  with erroneous temperature or pressure, or zero mixing ratio but non-zero flux, are removed. 8 indicates no values from 2013 and ozone concentration values used are from the slow sensor, the value for 23 m is linearly interpolated for measurements at 16.8 and 33.6 m; a indicates whether we made an assumption about air density in calculating  $v_d$  from ozone concentrations and fluxes received by the contact (in this case, we assume 25°C and 1013 hPa).



**Figure 1.** AM3DD evaluation of monthly daily (24-hour) mean ozone deposition velocity ( $v_d$ ) at sites with ozone eddy covariance fluxes (see Table 1) for sites with multiple years of data with at least a couple of months of data collected in a given year. Grey indicates the observational monthly average for a given year; black shows the multiyear average when available. Blue dashed lines show simulated  $v_d$  for the land cover type that best characterizes the site (blue text). For the observations, we calculate the monthly average  $v_d$  using a bootstrapping technique (see Clifton et al. (2017, 2019)). For a monthly average to be included, each hour of the day must have at least 25% data capture for the month. The error bars indicate the 95% confidence intervals.



**Figure 2.** Model evaluation of variability in ozone dry deposition with short-term observational data. (a)-(d) As in Figure 1 but for sites with short-term data. (e) Comparison of simulated and observationally based daily mean (24-hour) stomatal fractions of ozone dry deposition. Error bars on the observationally based values indicate two standard deviations across estimates given for a particular site and season; error bars on simulated values indicate two standard deviations across daily values. Black outlines on symbols represent sites where modeled LAI is less than  $1 \text{ m}^2 \text{ m}^{-2}$ , which may lead to underestimated stomatal fractions. Sites included are sites for which daily averages of the stomatal fraction were inferred from previous literature by Clifton et al. (2020).

414 We compile estimates of the stomatal fraction of ozone dry deposition over phys-  
415 iologically active vegetated landscapes from previous literature to evaluate simulated par-  
416 titioning between stomatal and nonstomatal deposition (Figure 2e). Estimates are based  
417 on ecosystem-scale ozone flux observations as well as micrometeorological observations  
418 used to infer stomatal uptake (*e.g.*, through inversions of water vapor fluxes or empir-  
419 ical stomatal conductance models) and resistances to turbulent and diffusive transport.  
420 We include here estimates that represent daily (24-hour) averages. While both the model  
421 and observationally based estimates show co-dominant roles for stomatal and nonstom-  
422 atal deposition, the simulated stomatal fraction is generally underestimated (only 37%  
423 of what it should be). However, sites with particularly low biases have very low simu-  
424 lated LAI (*e.g.*, 83% site-specific seasonal mean modeled stomatal fractions of  $<0.2$  have  
425  $<1 \text{ m}^2 \text{ m}^{-2}$  LAI), suggesting that the cause of the bias may be due to the model's in-  
426 ability to capture the amount of vegetation at these locations (to the extent that LAI  
427 is reported for the observational sites, most have higher LAI than this). Most sites lack  
428 the coincident measurements of LAI and stomatal fraction, which we need to directly  
429 evaluate the model's strength at capturing stomatal fractions where LAI is simulated ac-  
430 curately. Nonetheless, for all model grid cells with summer mean LAI  $>2 \text{ m}^2 \text{ m}^{-2}$  be-  
431 tween 35-50°N, the simulated summer stomatal fraction of ozone dry deposition is 0.4,  
432 matching the observationally based stomatal fraction (0.39). We therefore suggest that  
433 the model reasonably captures stomatal versus nonstomatal partitioning where substan-  
434 tial vegetation is simulated. In general, excessively low or high model LAI may imply  
435 a model overemphasis or underemphasis, respectively, of nonstomatal deposition.

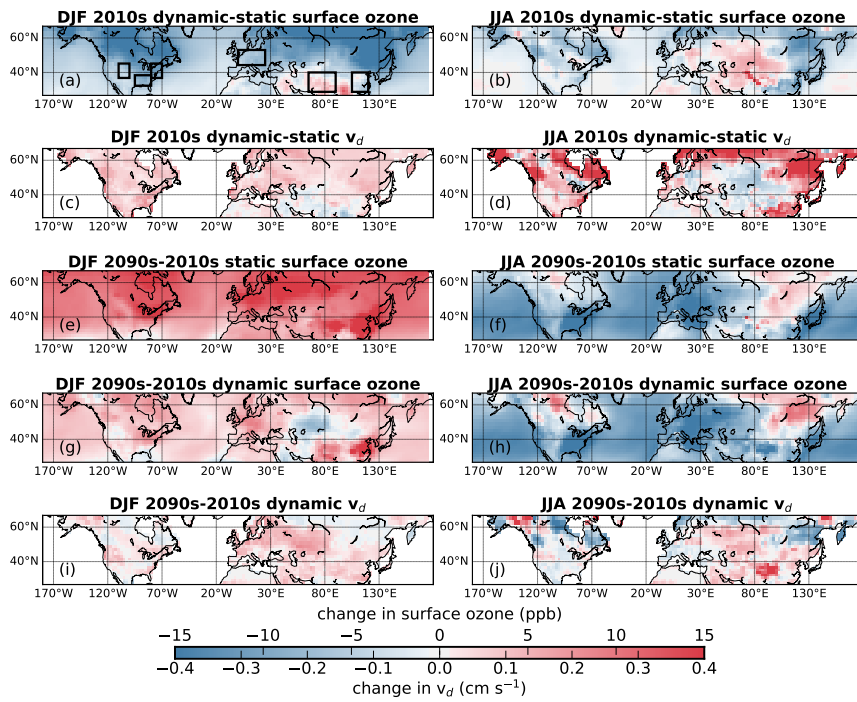
## 436 **4 Impact of dynamic ozone dry deposition scheme on present-day sur-** 437 **face ozone**

### 438 **4.1 Winter**

439 Winter surface ozone decreases by 10 ppb on average across northern mid-latitudes  
440 (40-55°N; land only) in response to higher (but still low) winter  $v_d$  in AM3DD versus  
441 AM3DD-staticO3DD (Figure 3a,c). For example, regional mean decreases for the regions  
442 outlined on Figure 3a (hereafter, highlighted regions) range from 3 to 10 ppb, except over  
443 central Asia where there are increases of 2 ppb. Winter  $v_d$  is 0.11 to 0.15  $\text{cm s}^{-1}$  in the  
444 monthly  $v_d$  climatology from GEOS-Chem for these regions, but 0.10 to 0.29  $\text{cm s}^{-1}$  as  
445 simulated by AM3DD.

446 Simulated winter surface ozone in AM3DD better matches most ground-based ob-  
447 servations across the northern hemisphere (Figure 4a,c,e), suggesting that ozone dry de-  
448 position may be key for representing winter surface ozone accurately. For the model eval-  
449 uation of surface ozone, we use 2008-2015 average daily mean mixing ratios from indi-  
450 vidual stations compiled for the Tropospheric Ozone Assessment Report (TOAR) (Schultz  
451 et al., 2017; Schultz et al., 2017). Over North America, Europe, and parts of Asia, the  
452 bias (simulated-observed) improvement is mostly within 1-15 ppb, but there are improve-  
453 ments of greater than 15 ppb at higher latitudes (*e.g.*, parts of Canada). At a couple of  
454 the most northern sites in Alaska and Scandinavia, surface ozone becomes too low in AM3DD.  
455 Over central Asia, the bias changes sign, but is small.

456 Reductions in winter surface ozone at any location may stem from local, upwind,  
457 and remote increases in ozone dry deposition. The winter ozone bias decreases by 5-12  
458 ppb in the lower troposphere in AM3DD relative to AM3DD-staticO3DD across north-  
459 ern mid-latitudes and boreal regions (40-65°N; land only) and remote locations where  
460 ozone sondes are regularly launched (Tilmes et al., 2012) (Figure 5) suggesting that ozone  
461 dry deposition influences baseline ozone, defined as ozone not recently influenced by lo-  
462 cal precursor emissions (HTAP, 2010).



**Figure 3.** Winter (December-February, or DJF) and summer (June-August, or JJA) differences between AM3DD (dynamic) and AM3DD-staticO3DD (static) for surface ozone mixing ratios and ozone deposition velocity ( $v_d$ ) at the 2010s, and differences between the 2090s and 2010s for  $v_d$  and surface ozone in AM3DD. We also show surface ozone differences between the 2090s and 2010s in AM3DD-staticO3DD. Black boxes on (a) represent regional definitions used in the paper and in subsequent figures.

463 Winter  $v_d$  is zero at northern latitudes in AM3DD-staticO3DD where there is snow,  
464 defined in GEOS-Chem as albedo  $>0.4$ . Winter  $v_d$  is only lower in AM3DD versus AM3DD-  
465 staticO3DD over parts of Asia (Figure 3c). Differences in  $v_d$  in these regions likely stem  
466 from slightly higher LAI in the satellite-based climatology used in GEOS-Chem (Fig-  
467 ure S1). At other mid-latitudes,  $v_d$  in AM3DD is higher than AM3DD-staticO3DD (*e.g.*,  
468 by 0.02 to 0.14  $\text{cm s}^{-1}$ ) and is almost completely dominated by ozone dry deposition to  
469 the ground (Figure 6a,c,e,g). Winter  $v_d$  in boreal regions with coniferous forests is dom-  
470 inated by uptake to cuticles (Figure 6a,c,e,g). While the comparison between LAI sim-  
471 ulated by the model and LAI in the satellite-based climatology used in GEOS-Chem sug-  
472 gests near-zero LAI in boreal forests during winter and thus an overestimate of LAI in  
473 AM3DD, satellite-based estimates of LAI over boreal regions are particularly uncertain  
474 due to snow contamination and low solar zenith angle (Fang et al., 2013, 2019).

475 Our parameterization addresses observational evidence that (1) ozone dry deposi-  
476 tion to snow-covered surfaces is low but nonzero (Helmig et al., 2007), (2) winter  $v_d$   
477 is lower over snow-covered versus bare surfaces in temperate regions (Padro et al., 1992;  
478 Stocker et al., 1995; Helmig et al., 2007; Matichuk et al., 2017), and (3) ozone dry de-  
479 position to snow-covered forests is relatively high compared to other snow-covered sur-  
480 faces (Z. Wu et al., 2016; Neiryck & Verstraeten, 2018). While there is uncertainty in  
481 the initial resistances and other parameters employed here, as well as the exact processes  
482 controlling winter ozone dry deposition, our results suggest that considering this evidence  
483 and a more dynamic representation of snow cover may be important for capturing tro-  
484 pospheric ozone abundances accurately.

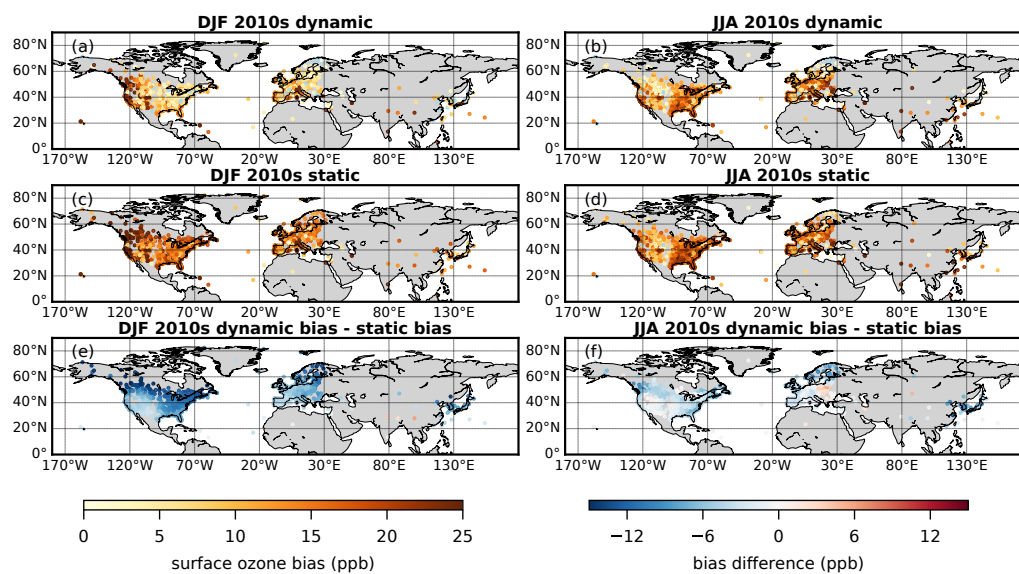
## 485 4.2 Summer

486 During June-August, surface ozone decreases on average by 5 ppb in AM3DD re-  
487 lative to AM3DD-staticO3DD over boreal latitudes where there is higher  $v_d$  in AM3DD  
488 (Figure 3b,d). Higher  $v_d$  over boreal latitudes is due to high stomatal and cuticular de-  
489 position to boreal coniferous forests (Figure 6i,k,m). The summer surface ozone bias re-  
490 duces by 1-10 ppb at all boreal monitoring sites except one (Figure 4b,d,f). However,  
491 LAI over much of the boreal forested region is higher than a satellite-based climatology  
492 (Figure S2), suggesting that ozone dry deposition may be too high over boreal forests  
493 and thus the substantial decrease in boreal surface ozone overestimated.

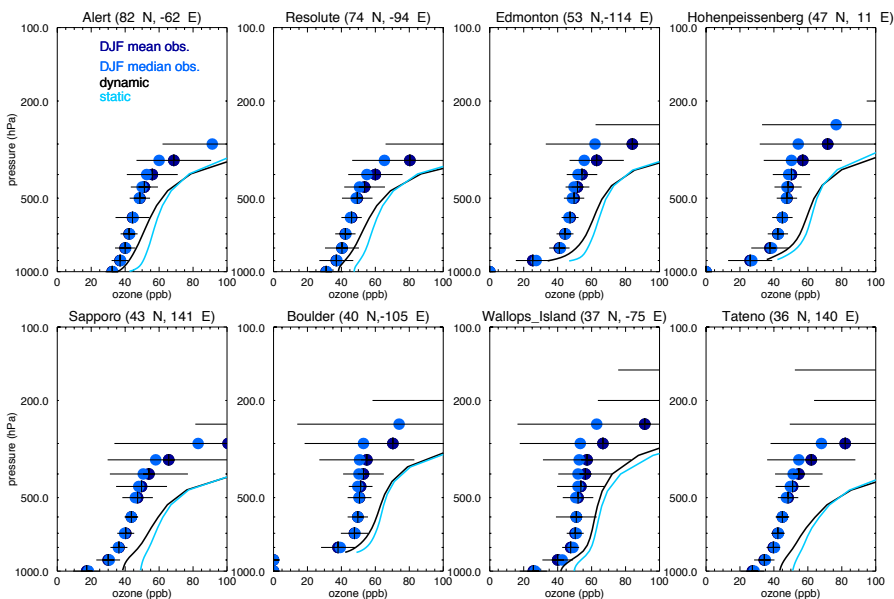
494 Over mid-latitudes, the sign of the change in summer surface ozone with dynamic  
495 ozone dry deposition varies (Figure 3b). Dynamic ozone dry deposition decreases the sum-  
496 mer mean surface ozone bias over North America and Europe by 2-7 ppb, with the ex-  
497 ceptions of eastern Europe and parts of the Great Lakes region of the US and western  
498 US where dynamic ozone dry deposition exacerbates the bias by 1-5 ppb (Figure 4f). Dy-  
499 namic ozone dry deposition decreases the summer mean ozone bias over east Asia by up  
500 to 10 ppb, but worsens the bias at the limited monitoring sites in other parts of Asia.  
501 Model LAI overestimates in south China may suggest a  $v_d$  overestimate there, but ozone  
502 flux measurements are needed to confirm this. In general, the ozone bias is worse in re-  
503 gions where simulated LAI is lower than the satellite-based estimate (Figure S2), sug-  
504 gesting that  $v_d$  is underestimated because there is not enough vegetation. Due to the  
505 short summer surface ozone lifetime (*e.g.* a few days over continental northern mid-latitude  
506 regions), surface ozone differences between AM3DD and AM3DD-staticO3DD tend to  
507 mirror  $v_d$  differences (Figure 3b,d).

508 Summer mean decreases up to 7 ppb in surface ozone occur over the southeast (SE)  
509 US. Such decreases may at least in part be due to wet cuticular deposition in AM3DD  
510 (Figure 6k), which is not simulated by the Wesely scheme in GEOS-Chem. The lack of  
511 wet cuticular deposition in most deposition schemes may thus contribute to the positive  
512 bias in modeled SE U.S. surface ozone (Fiore et al., 2009; Travis et al., 2016). Travis and





**Figure 4.** Winter (December-February, or DJF) and summer (June-August, or JJA) model evaluation using 2008-2015 mean surface ozone mixing ratios from individual stations compiled and calculated for the Tropospheric Ozone Assessment Report (TOAR) (Schultz et al., 2017; Schultz et al., 2017). In (a)-(d), we show the surface ozone bias (simulated minus observed) at each site for AM3DD (dynamic) and AM3DD-staticO3DD (static). Negative biases are shown in light blue. In (e)-(f), we show the difference in the biases. Negative values indicate improvement. If the bias is negative under AM3DD-staticO3DD then the site is not shown on (e)-(f) (the few removed sites are shown in light blue on panels (c)-(d)). We remove sites with less than 50% hourly data coverage (averaged over all winter or summer days in 2008 to 2015) and less than 50% of yearly coverage. We also discard sites characterized as traffic, industry, urban, and suburban by individual monitoring networks in order to lessen the influence of polluted urban air on our coarse-scale model evaluation, with the caveat that most sites are not classified.



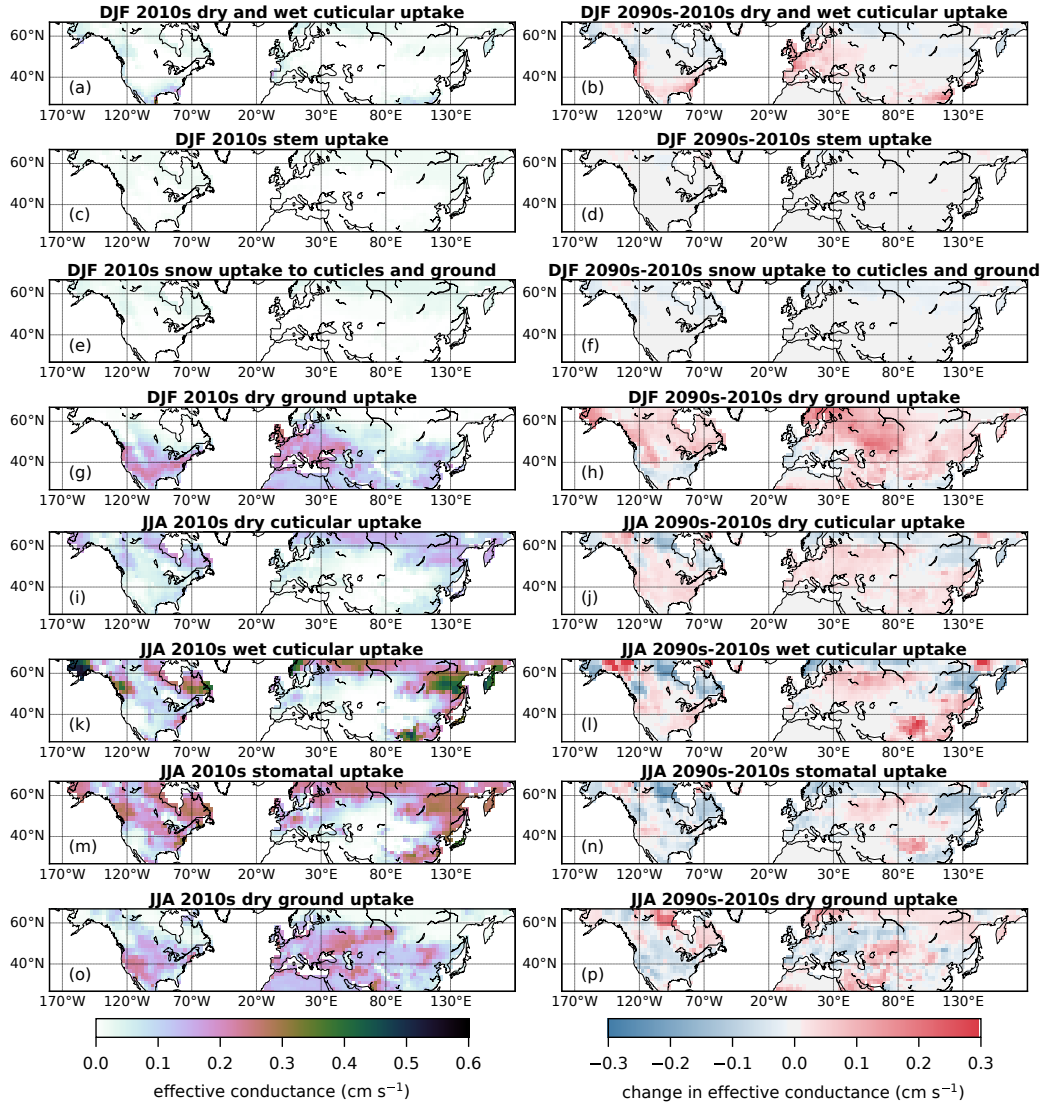
**Figure 5.** Winter (December-February, or DJF) model evaluation using 1995-2009 ozone vertical profiles from ozone sonde observations at individual stations north of  $35^{\circ}\text{N}$  from Tilmes et al. (2012) for AM3DD (dynamic) and AM3DD-staticO3DD (static).

513 Jacob (2019) also suggest the absence of this process prevents GEOS-Chem from cap-  
 514 turing low SE U.S. surface ozone on rainy days.

515 Besides summer mean differences in surface ozone between AM3DD and AM3DD-  
 516 staticO3DD, there are also differences in daily probability distributions (Figure 7a-f).  
 517 For the SE US, the distribution decreases and there are larger changes for wet days ( $>6$   
 518  $\text{mm day}^{-1}$  precipitation as defined in Travis and Jacob (2019)) versus all days in AM3DD  
 519 relative to AM3DD-staticO3DD, suggesting that high  $v_d$  on rainy days drives regional  
 520 surface ozone decreases with dynamic ozone dry deposition. For the Northeast (NE) US  
 521 and the InterMountain West (IMW) US, the mode of the distribution decreases, and the  
 522 distribution shifts towards lower values. For central Asia, the mode of the distribution  
 523 also decreases but the distribution shifts towards higher values. For central Europe, the  
 524 distribution widens, with higher and lower surface ozone extremes.

525 Daily variability in  $v_d$  in AM3DD may drive the changes in the distribution of sur-  
 526 face ozone across days. However, there is some evidence that mean changes in  $v_d$  may  
 527 contribute to changes in relative variability in surface ozone. For example, reducing  $v_d$   
 528 by 35% over drought-stricken regions of the eastern US in 1988 with the version of AM3  
 529 employing the monthly  $v_d$  climatology shifts the ozone distribution towards higher val-  
 530 ues, but also slightly decreases the mode of the distribution (M. Lin et al., 2017), im-  
 531 plying a nonlinear ozone response to a mean shift in  $v_d$ . Disentangling contributions to  
 532 the changes in the surface ozone distribution from daily-varying  $v_d$  versus a nonlinear  
 533 ozone sensitivity to  $v_d$  is not possible with our simulations. Nonetheless, strong corre-  
 534 lations between  $v_d$  and surface ozone on daily timescales (Figure 7g) suggest that day-  
 535 to-day variability in ozone dry deposition plays an important role in shaping the surface  
 536 ozone distribution across days.

537 Kavassalis and Murphy (2017) hypothesize variability in stomatal ozone dry dep-  
 538 osition influences daily variability in ozone pollution on the basis of the strong cor-  
 539 relation between observed surface ozone concentrations and vapor pressure deficit and a



**Figure 6.** Winter (December-January, or DJF) and summer (June-August, or JJA) effective conductances at the 2010s, and differences between the 2090s and 2010s under AM3DD. For a given season, we only show deposition pathways that substantially contribute to ozone deposition velocity ( $v_d$ ); the effective conductances shown sum to  $v_d$ . The change in the effective conductances sum to the net change in  $v_d$  from the 2010s to 2090s shown in Figure 3. For all panels, grid cells with less than 50% land are not included.

540 strong dependence of stomatal conductance on dryness. In AM3DD, nonstomatal de-  
541 position is an important fraction of the total ozone dry deposition (Figure 6i,k,m,o) and  
542 a key driver of daily variability in summer  $v_d$  (Figure 7i-l), suggesting that dynamic non-  
543 stomatal deposition also influences daily variability in surface ozone. In particular, wet  
544 cuticular and ground deposition vary, reflecting the influence of soil and leaf wetness, re-  
545 spectively, as well as in-canopy turbulence for the latter, and dominate the variability  
546 in  $v_d$  in many regions (Figure 7i-l).

547 The correlation between wet cuticular and stomatal deposition (Figure 7h) and the  
548 substantial magnitude and variability that each of these terms provides summer  $v_d$  (Fig-  
549 ures 6i,k,m,o and 7i-l) imply that an unambiguous attribution of increases in ozone pol-  
550 lution during drought to reductions in stomatal deposition may be challenging. M. Lin  
551 et al. (2019) use a similar version of the GFDL model to conclude that variations in stom-  
552 atal deposition drive variations in ozone pollution with drought. However, M. Lin et al.  
553 (2019) do not consider how variations in cuticular uptake with precipitation influence  
554 variability in  $v_d$  and thus their conclusion may need to be re-visited.

555 Most studies examining observed  $v_d$  after rain and dew report increases (Clifton  
556 et al., 2020). While laboratory and field chamber evidence support increases in cutic-  
557 ular deposition to wet leaves (Fuentes & Gillespie, 1992; Pleijel et al., 1995; Sun et al.,  
558 2016; Potier et al., 2017), whether increases in ecosystem-scale  $v_d$  after rain and dew are  
559 due to wet cuticular deposition is uncertain. For example, changes in other processes (*e.g.*,  
560 stomatal conductance, in-canopy chemistry) may contribute to observed increases (Altimir  
561 et al., 2006; Turnipseed et al., 2009; Clifton et al., 2019). Canopy interception of water  
562 is also an uncertain component of land models (Bonan & Levis, 2006; De Kauwe et al.,  
563 2013; Lian et al., 2018; Fan et al., 2019) and contributes to uncertainty in simulated wet  
564 cuticular deposition. The amount of canopy-intercepted precipitation in LM3 is lower  
565 than observation-based estimates (Milly et al., 2014) and additional uncertainty includes  
566 the duration and fraction of wet leaves.

567 In general, AM3DD may not capture the partitioning of  $v_d$  to individual pathways  
568 accurately due to process and parameter uncertainty (*e.g.*,  $m$ ,  $d_0$ , all initial resistances).  
569 Indeed, recent work identifies factor of 2-3 differences in simulated  $v_d$  due to process rep-  
570 resentation and parameter choice (Z. Wu et al., 2018; Wong et al., 2019). Given that AM3DD  
571 seems to capture the magnitude of  $v_d$ , model LAI under- or overestimates (Figure S2)  
572 may imply a nonstomatal deposition over- or underemphasis, respectively. Comparisons  
573 with other models that prognostically simulate the components of ozone dry deposition  
574 (*i.e.*, LAI, soil moisture) will be useful for assessing confidence in the contribution of dif-  
575 ferent processes to ozone dry deposition as represented in current models.

## 576 **5 21<sup>st</sup>-century changes in surface ozone from dynamic ozone dry de-** 577 **position**

### 578 **5.1 Winter northern mid-latitudes**

579 Over northern mid-latitudes, winter surface ozone increases with the 21<sup>st</sup>-century  
580 reductions in anthropogenic nitrogen oxide ( $\text{NO}_x$ ) emissions (*i.e.*, 2010-to-2090 decreases  
581 of 57-69% for the highlighted regions) and doubling of global methane under RCP8.5 (*i.e.*,  
582 105% increase from 2010 to 2090) (Gao et al., 2013; Clifton et al., 2014). More specifi-  
583 cally, reductions in regional  $\text{NO}_x$  emissions under RCP8.5 over polluted mid-latitudes  
584 lead to a reversal of surface ozone seasonality from a summer to a winter peak and the  
585 global methane doubling amplifies hemispheric-scale ozone (Clifton et al., 2014).

586 We find here that increasing winter  $v_d$  during the 21<sup>st</sup> century tempers the rise in  
587 winter surface ozone over mid-latitudes in AM3DD relative to AM3DD-staticO3DD (Fig-  
588 ure 3e,g,i). For example, 21<sup>st</sup>-century increases in winter surface ozone are lower on av-  
589 erage by 4-8 ppb in AM3DD relative to AM3DD-staticO3DD for highlighted regions. Over

590 some parts of Asia, changes in local and remote ozone dry deposition tip the balance to-  
591 wards 21<sup>st</sup>-century decreases in winter ozone.

592 Higher winter  $v_d$  by the 2090s at mid-latitudes mainly reflects higher ground de-  
593 position and higher dry and wet cuticular deposition (Figure 6b,d,f,h). There is higher  
594 ground deposition due to less snow. Andersson and Engardt (2010) also find that de-  
595 creasing snow over Europe with climate warming is an important driver of regional  $v_d$   
596 and ozone pollution for their April-October analysis. Increases in winter  $v_d$  from higher  
597 cuticular deposition are likely associated with warmer winters and higher LAI (Figure  
598 S3) from the long-term effects of carbon dioxide fertilization (*i.e.*, plants accumulate more  
599 biomass under high carbon dioxide).

## 600 5.2 Summer northern mid-latitudes

601 Large summer decreases in surface ozone from the 2010s to the 2090s over polluted  
602 northern mid-latitudes occur as regional anthropogenic  $\text{NO}_x$  emissions decline under RCP8.5  
603 (Gao et al., 2013; Clifton et al., 2014; Rieder et al., 2018). Similar to AM3DD-staticO3DD,  
604 summer surface ozone decreases over most mid-latitudes in AM3DD (Figure 3f,h). For  
605 highlighted regions, the 21<sup>st</sup>-century decrease in surface ozone is -7 to -17 ppb in AM3DD  
606 versus -2 to -19 ppb in AM3DD-staticO3DD; the decrease weakens by about 1 ppb in  
607 AM3DD except over central and east Asia where the decreases are the same or become  
608 stronger by 4 ppb, respectively.

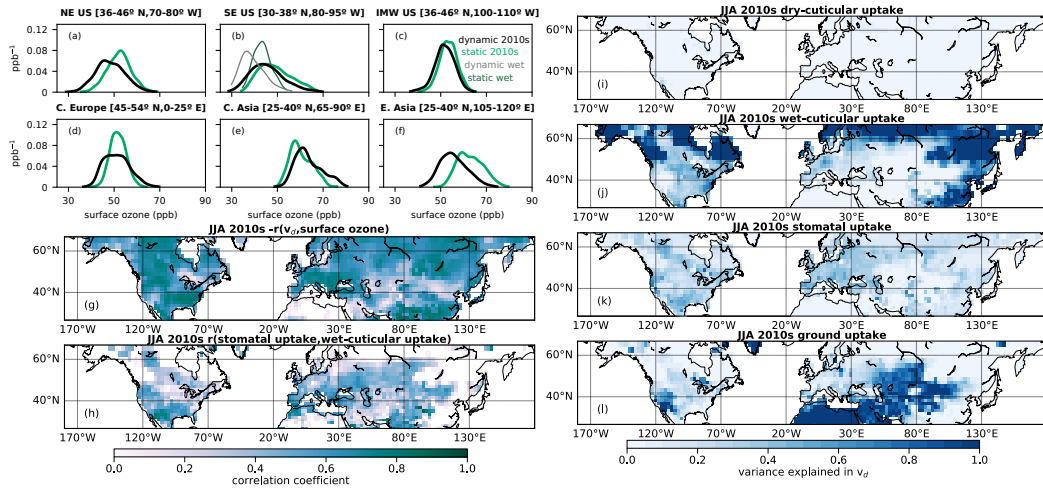
609 Over several mid-latitude regions, opposing changes in individual deposition path-  
610 ways from the 2010s to the 2090s offset each other, leading to little net 21<sup>st</sup>-century change  
611 in  $v_d$ . For example, summer dry cuticular deposition increases nearly everywhere from  
612 the long-term effects of carbon dioxide fertilization promoting leaf biomass accumula-  
613 tion (Figure 6j). Wet cuticular deposition increases or does not change at most mid-latitudes  
614 (Figure 6l); regions with increases in wet cuticular deposition are regions with increases  
615 in rainfall and regions with no change are regions with decreases in rainfall (Figure S4b).  
616 Ground deposition decreases or does not change in most mid-latitude regions, except west-  
617 ern Asia (Figure 6h). Changes in ground deposition mostly reflect higher LAI, which raises  
618 the resistance to in-canopy turbulence and decreases ground uptake, rather than changes  
619 in soil wetness, which are mostly decreases and would lead to increases in ground up-  
620 take (Figures S4a,d). Summer stomatal deposition either does not change or decreases  
621 over most mid-latitude regions (Figure 6n) despite widespread increases in LAI, likely  
622 due to increased dryness and the short-term (*i.e.*, instantaneous) effects of carbon diox-  
623 ide that decrease stomatal conductance (Figure S4c,d). Exceptions include western Asia  
624 and the western US where there is vegetation at end of the century but not at the be-  
625 ginning (compare Figures S2b and S4a).

## 626 5.3 Summer and winter boreal regions

627 With the prescription of land use change under RCP8.5 and the expansion of de-  
628 ciduous forests into boreal latitudes simulated by the vegetation dynamics in LM3, there  
629 are 21<sup>st</sup>-century decreases in winter and summer cuticular deposition (Figures 6b,f,j,l)  
630 over boreal regions with conifers at the 2010s. Such decreases likely occur because the  
631 model generally simulates lower LAI for deciduous forests, pastures, and crops relative  
632 to coniferous forests (not shown). There are 21<sup>st</sup>-century decreases in summer stomatal  
633 deposition over these boreal regions (Figure 6n), likely following decreases in LAI but  
634 also the short-term impact of high carbon dioxide. In the regions north of 50°N with de-  
635 ciduous forests throughout the 21<sup>st</sup> century, increases in winter and summer  $v_d$  follow  
636 less snow (winter only) and higher LAI from carbon dioxide fertilization.

637 Our findings contrast with S. Wu et al. (2012) who find widespread increases in bo-  
638 real summer  $v_d$  between 2000 and 2100. Differences in  $v_d$  between AM3DD and their





**Figure 7.** Daily variability in surface ozone and ozone dry deposition. (a)-(f) Summer (June-August) probability density functions of daily regional average surface ozone mixing ratios for the 2010s in several northern mid-latitude regions for AM3DD (dynamic) and AM3DD-staticO3DD (static) estimated with a Gaussian kernel density. The regions are indicated with black lines on Figure 3a. For the southeast US, we also include probability density functions for wet days only (defined as 6 mm day<sup>-1</sup> on a regional average basis). (g) Correlation coefficient between day-to-day variability in summer surface ozone and ozone deposition velocity ( $v_d$ ) in AM3DD. (h) Correlation coefficient between day-to-day variability in summer effective stomatal and wet-cuticular conductances in AM3DD. For (g)-(h), white space on land denotes correlations outside the color bar. In (g), all correlations shown are negative but displayed as positive. (i)-(l) Variance explained in summer daily  $v_d$  by individual deposition pathways for AM3DD. We use the variance formula for variables that are not independent from each other ( $Var(\sum_{i=1}^n X_i) = \sum_{i=1}^n Var(X_i) + 2 \sum_{1 \leq i < j < n} Cov(X_i, X_j)$ ) because each effective conductance is the fraction of deposition through a certain pathway multiplied by  $v_d$ . For all panels, grid cells with less than 50% land are not included.

639 model (the dynamic vegetation model described in Sitch et al. (2003)) result from dif-  
640 ferent prognostically determined LAI (*i.e.*, their model shows 21<sup>st</sup>-century LAI increases  
641 over boreal regions), prescriptions of land use change, and stomatal conductance param-  
642 eterizations. S. Wu et al. (2012) use a Jarvis (1976) stomatal conductance model rather  
643 than a coupled net photosynthesis-stomatal conductance model as used here. While their  
644 stomatal conductance parameterization considers the long-term effect of carbon diox-  
645 ide on LAI, it does not consider the short-term effect on stomatal conductance.

646 In general, 21<sup>st</sup>-century carbon dioxide fertilization is uncertain (Wieder et al., 2015;  
647 N. G. Smith et al., 2016; W. K. Smith et al., 2016; Yuan et al., 2019; Terrer et al., 2016;  
648 Sulman et al., 2019; Humphrey et al., 2018; Green et al., 2019; Friedlingstein et al., 2006;  
649 Gerber et al., 2010). For example, changes in other processes may offset or exacerbate  
650 the impacts of high carbon dioxide on stomatal conductance and LAI (*e.g.*, nutrient lim-  
651 itation). A better understanding of carbon dioxide fertilization will not only lead to more  
652 accurate projections of stomatal deposition, but also nonstomatal deposition.

## 653 6 Conclusion

654 Limited representation of ozone dry deposition in atmospheric chemistry models  
655 hampers understanding of ozone pollution because simulated surface ozone is sensitive  
656 to  $v_d$  (J.-T. Lin et al., 2008; Walker, 2014; Hogrefe et al., 2018). Here we use a new ver-  
657 sion of the NOAA GFDL global chemistry-climate model, AM3DD, that leverages the  
658 dynamics of the underlying land model to simulate dry deposition of some aerosols and  
659 reactive trace gases, including ozone. Particularly novel features of the dynamic ozone  
660 dry deposition scheme are dependencies of nonstomatal deposition processes on soil mois-  
661 ture, canopy humidity, and canopy interception of water and snow and stomatal depo-  
662 sition on photosynthesis, vapor pressure deficit, and soil moisture. We use this new tool  
663 to investigate the influence of ozone dry deposition on surface ozone at northern mid-  
664 latitudes at the beginning and end of the 21<sup>st</sup> century. While stomatal deposition has  
665 long been recognized as an important driver of ozone dry deposition, we show that the  
666  $v_d$  spatial distribution, daily variability, and 21<sup>st</sup>-century changes also depend on non-  
667 stomatal deposition.

668 The new version of the GFDL model improves the simulation of winter ozone at  
669 surface monitoring sites and in the lower troposphere at remote sites relative to the ver-  
670 sion of the model driven with a  $v_d$  climatology. Higher simulated winter  $v_d$  in AM3DD  
671 reflects our use of interactive snow dynamics and recognizing non-negligible winter ozone  
672 dry deposition, as supported by observations. A major finding from our study is that  
673 winter ozone dry deposition influences baseline ozone, suggesting that remote ozone dry  
674 deposition is an important lever on a given region's ozone pollution. We also find that  
675 large-scale increases in winter  $v_d$  during the 21<sup>st</sup> century under RCP8.5 limit the influ-  
676 ence of rising global methane on surface ozone (*e.g.*, Clifton et al. (2014)). For exam-  
677 ple, the change in winter surface ozone from the 2010s to 2090s with dynamic ozone dry  
678 deposition is 1 to 13 ppb over the northern mid-latitude regions highlighted here versus  
679 6 to 21 ppb with the climatology.

680 The dynamic ozone dry deposition scheme generally leads to -4 to +7 ppb changes  
681 in mean summer surface ozone at the 2010s over northern mid-latitudes relative to the  
682 simulation forced with a  $v_d$  climatology. We find that daily variations in summer  $v_d$  with  
683 meteorology and biophysics, including from nonstomatal deposition processes, contribute  
684 to daily variations in ozone pollution. Evidence includes differences in daily ozone prob-  
685 ability distributions between simulations with dynamic ozone dry deposition versus the  
686 climatology, daily correlations between surface ozone and  $v_d$  in the dynamic simulation,  
687 and the high fraction of variance explained by nonstomatal deposition in simulated daily  
688 variations in  $v_d$ . Our new dry deposition configuration supports a role for ozone dry de-  
689 position on rainy days in the pervasive summer surface ozone bias over the southeast US.

In general, simulated cuticular deposition varies similarly to stomatal deposition, suggesting unambiguous attribution of variations in  $v_d$  and ozone pollution to stomatal deposition may be challenging. Studies pinpointing the drivers of day-to-day variability in observed  $v_d$  will be useful for ensuring that regional-to-global models capture the response of summer ozone dry deposition to meteorological and biophysical variability accurately.

Mostly 21<sup>st</sup>-century changes in summer surface ozone at northern mid-latitudes under RCP8.5 are similar with dynamic ozone dry deposition (around 1 ppb difference). One exception is east Asia where increasing  $v_d$  leads to a 4 ppb stronger decrease in summer surface ozone. In general, there are changes in summer ozone deposition pathways with changes in rainfall, dryness and carbon dioxide. However, changes in individual pathways tend to offset one another and thus there is not much impact on the change in summer surface ozone. The extent to which this offsetting occurs, however, depends fundamentally on assumptions inherent to the representation of different depositional processes in the model. Given the reliance of all ozone dry deposition parameterizations on myriad uncertain tuning parameters that determine the magnitude of the 21<sup>st</sup>-century changes in individual deposition processes, improved understanding of such processes is needed (*e.g.*, Clifton et al. (2020)).

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Figure 1.

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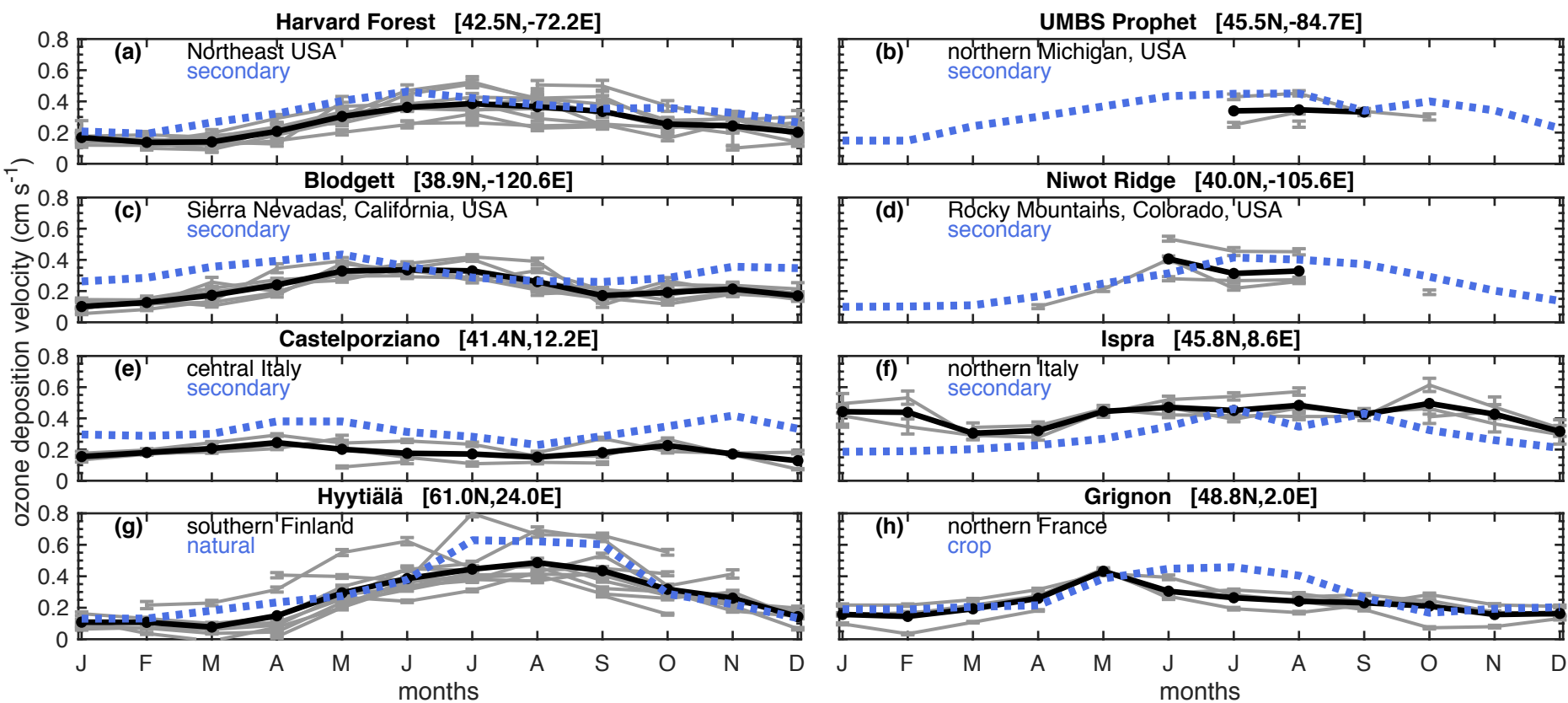




Figure 2.

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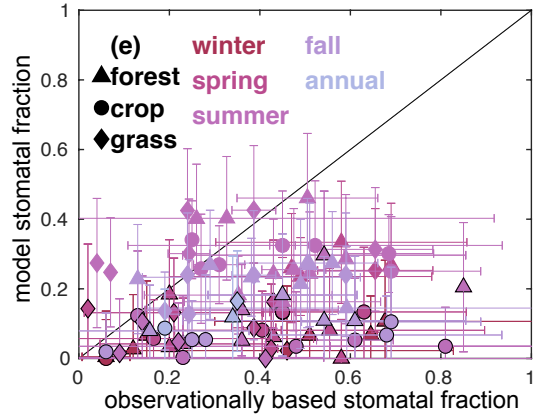
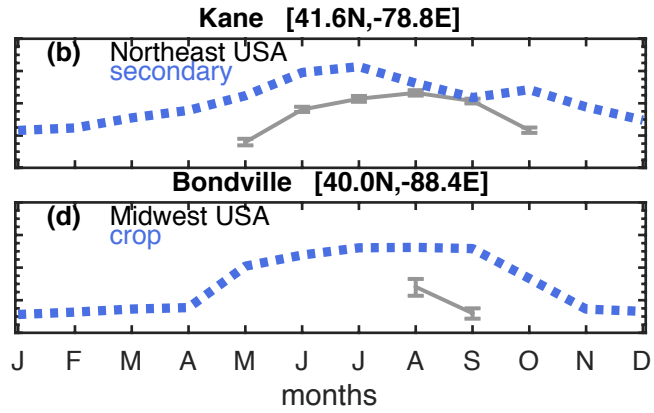
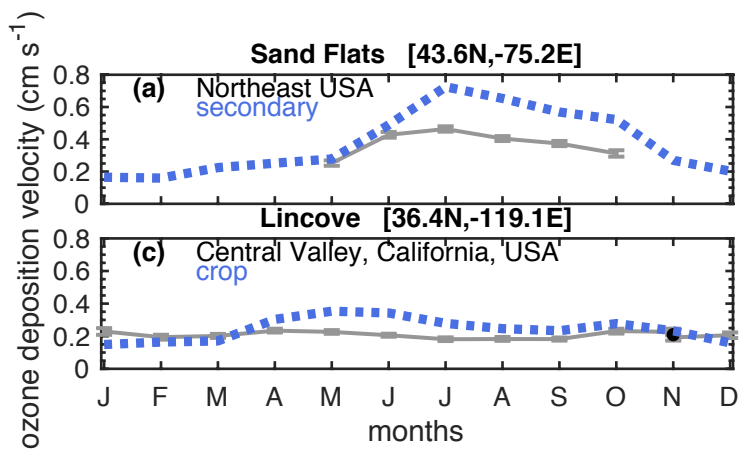


Figure 3.

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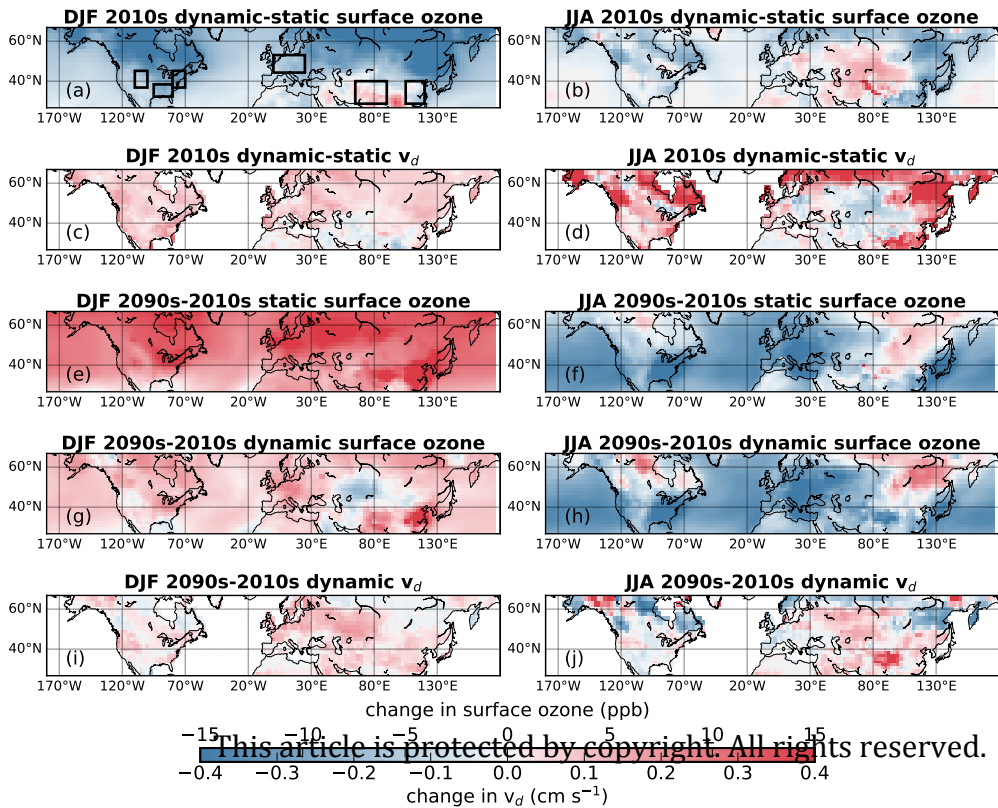
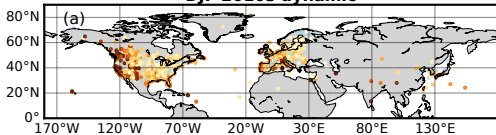


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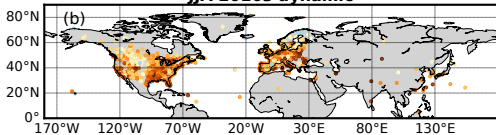
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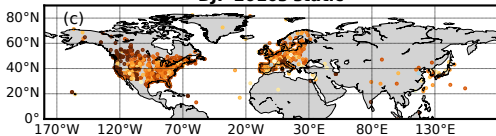
**DJF 2010s dynamic**



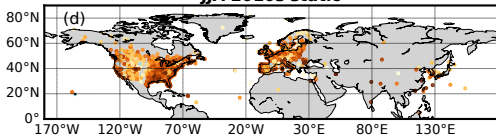
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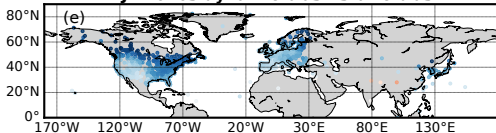
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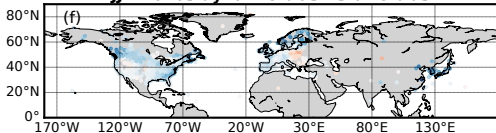
**JJA 2010s static**



**DJF 2010s dynamic bias - static bias**



**JJA 2010s dynamic bias - static bias**



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Figure 5.

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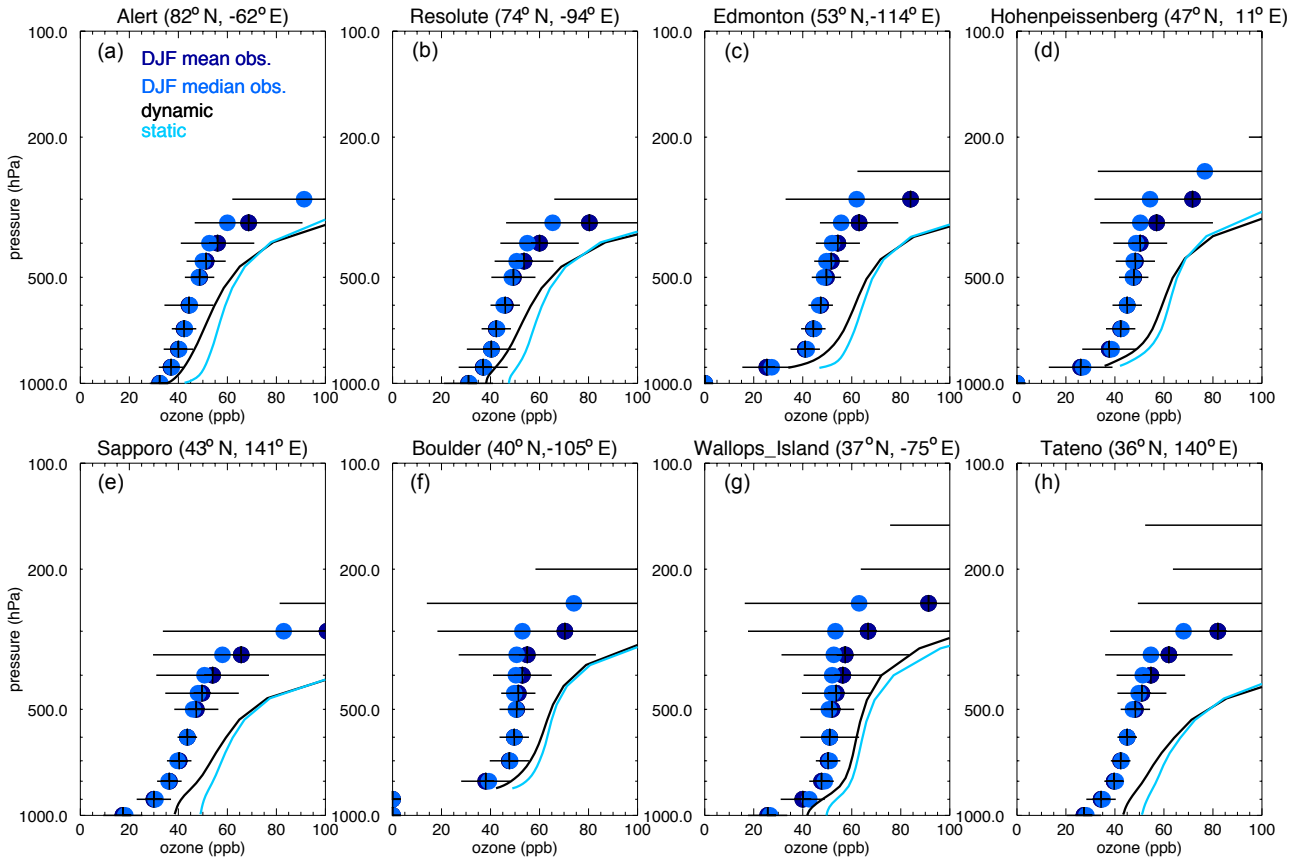


Figure 6.

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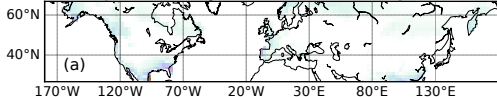
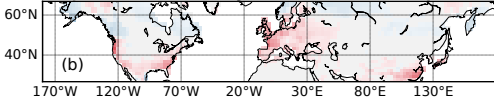
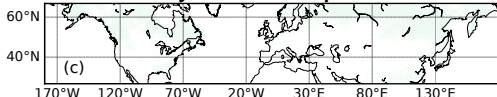
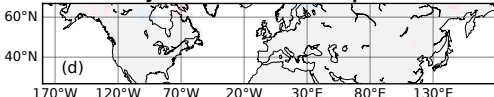
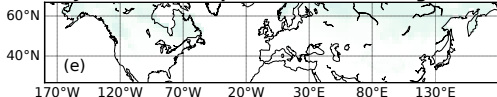
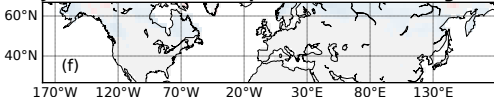
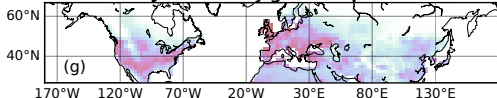
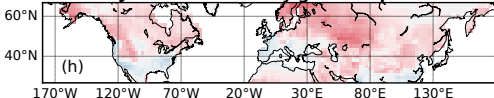
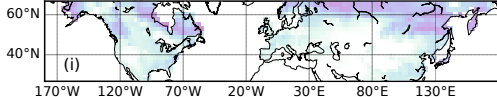
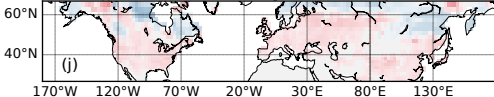
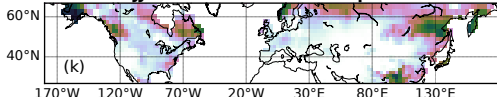
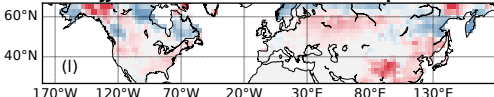
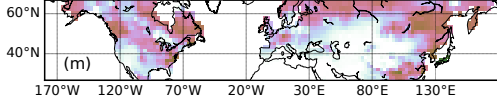
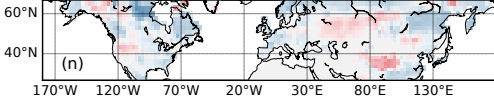
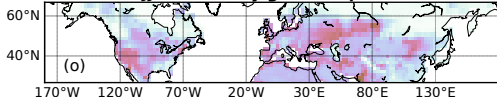
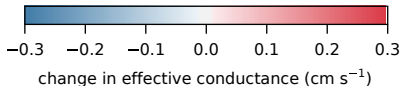
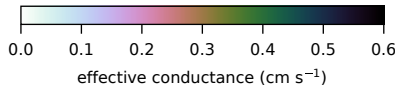
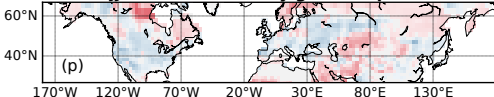
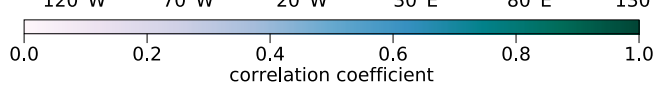
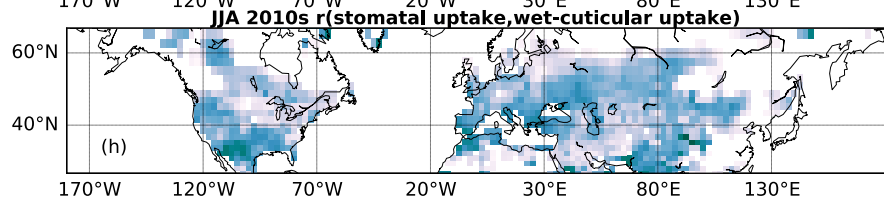
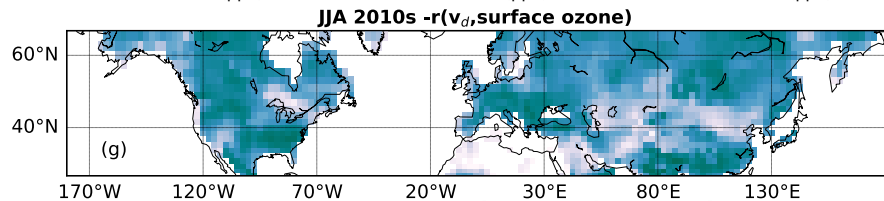
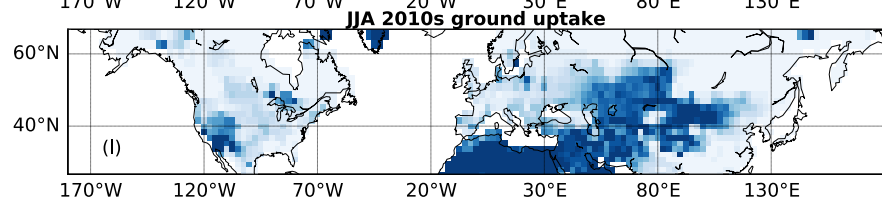
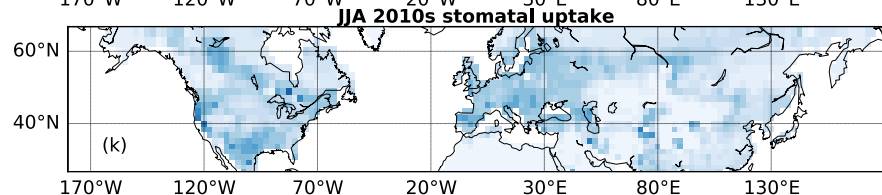
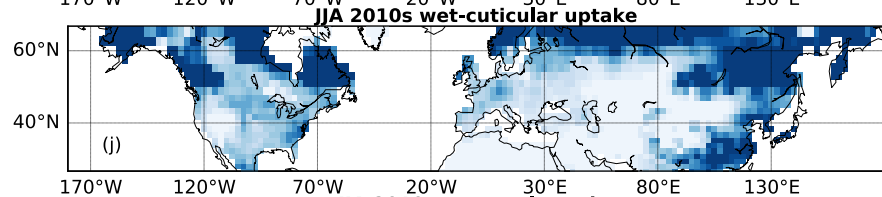
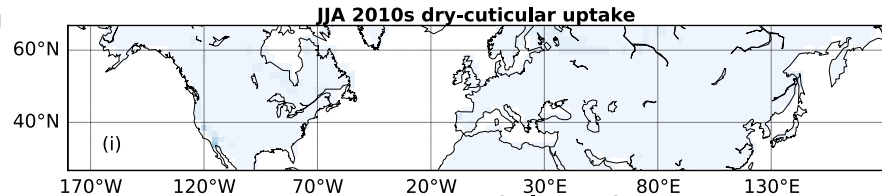
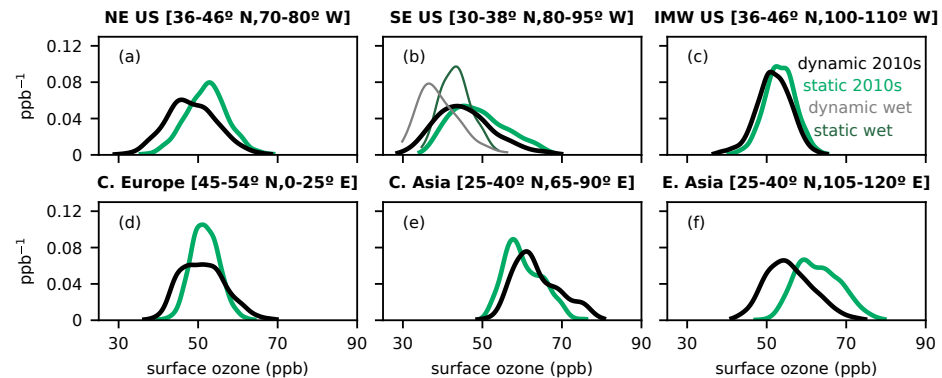
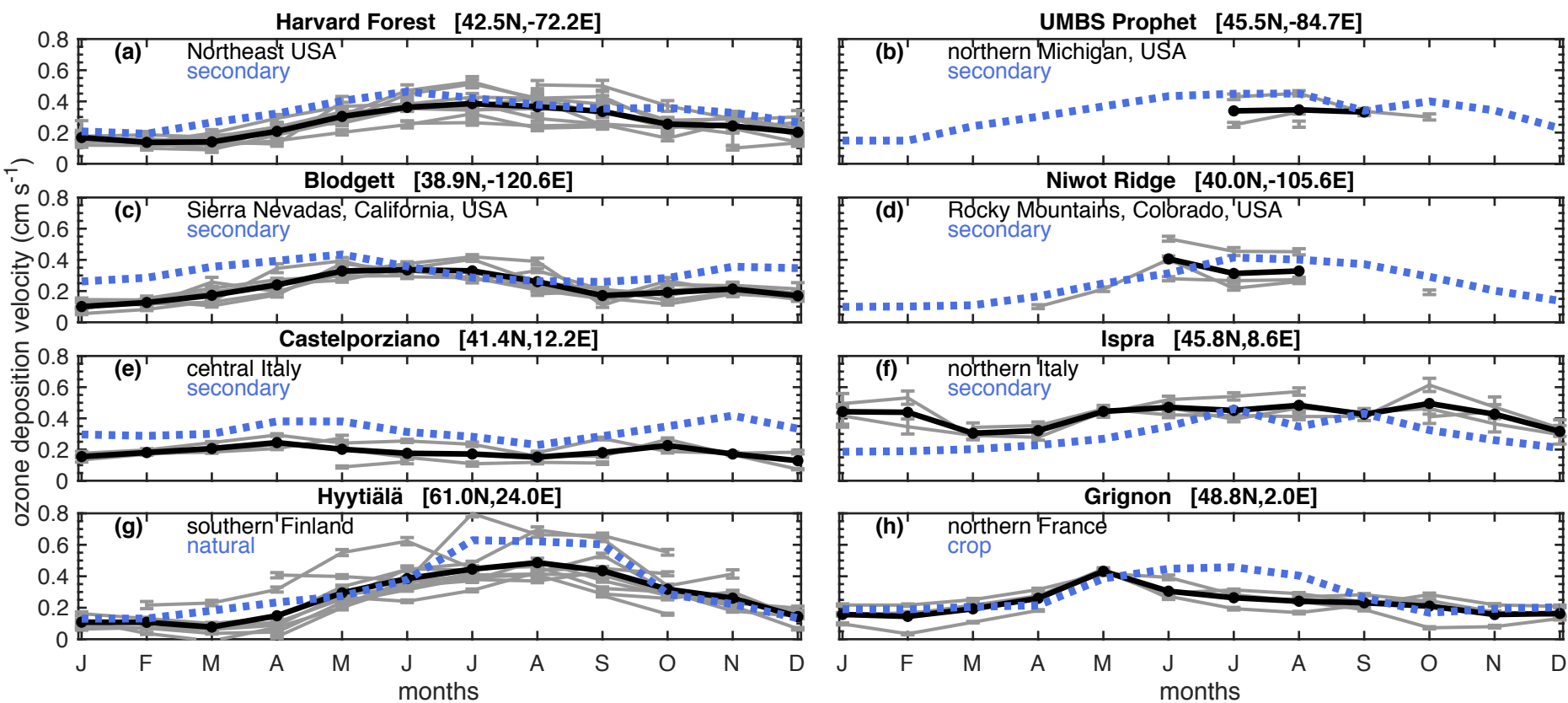
**DJF 2010s dry and wet cuticular uptake****DJF 2010s-2010s dry and wet cuticular uptake****DJF 2010s stem uptake****DJF 2010s-2010s stem uptake****DJF 2010s snow uptake to cuticles and ground****DJF 2010s-2010s snow uptake to cuticles and ground****DJF 2010s dry ground uptake****DJF 2010s-2010s dry ground uptake****JJA 2010s dry cuticular uptake****JJA 2010s-2010s dry cuticular uptake****JJA 2010s wet cuticular uptake****JJA 2010s-2010s wet cuticular uptake****JJA 2010s stomatal uptake****JJA 2010s-2010s stomatal uptake****JJA 2010s dry ground uptake****JJA 2010s-2010s dry ground uptake**

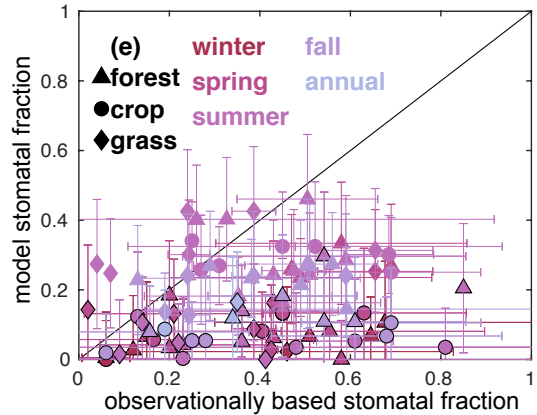
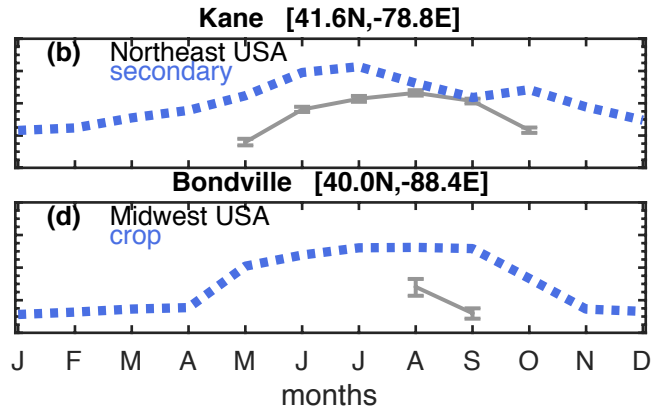
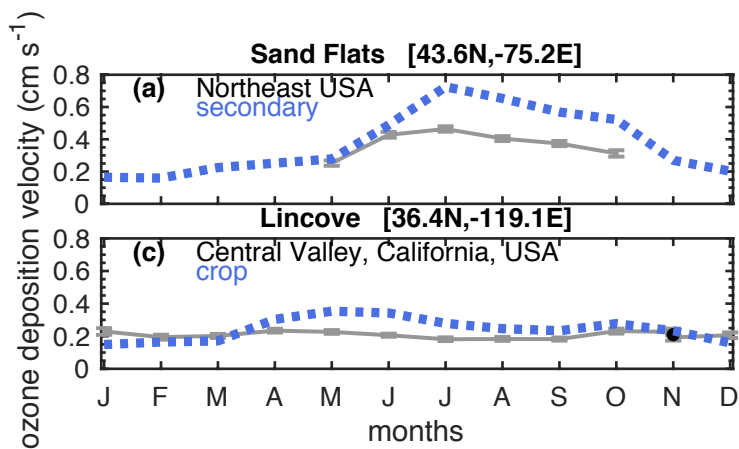


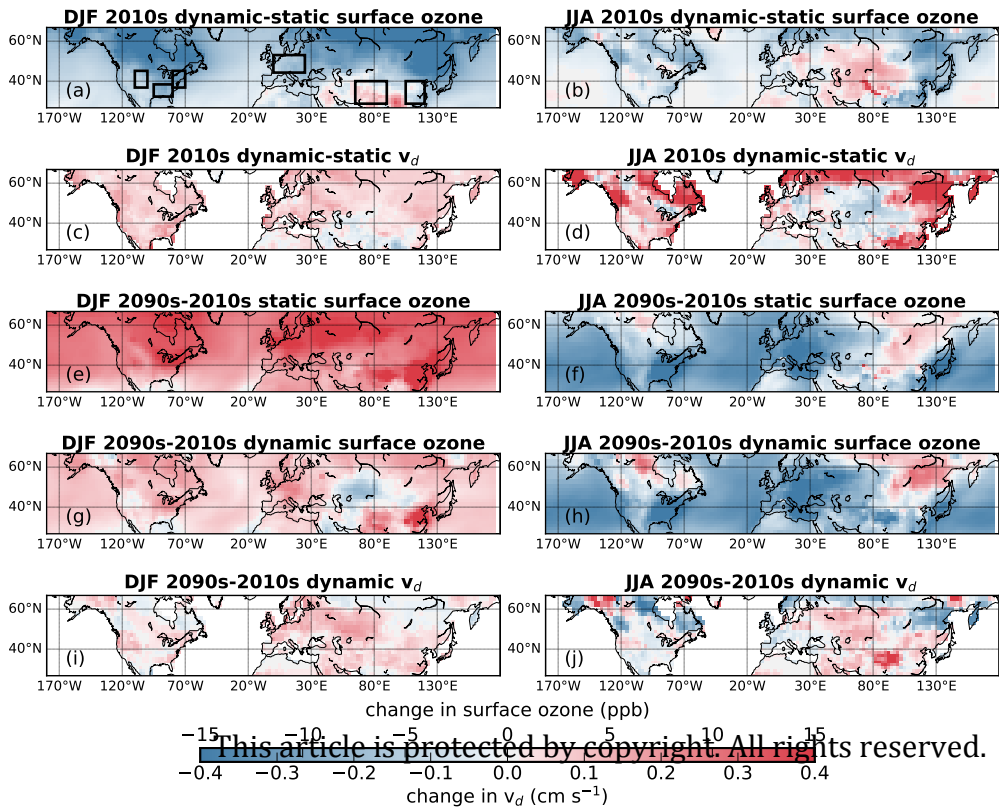
Figure 7.

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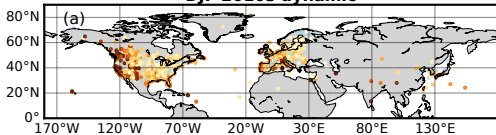




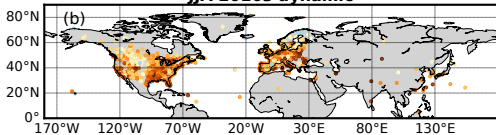




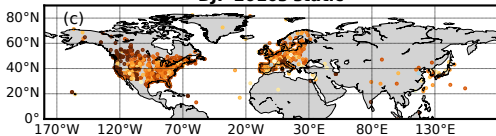
**DJF 2010s dynamic**



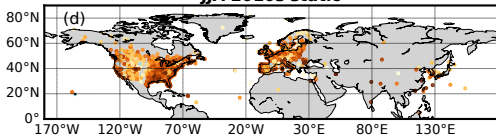
**JJA 2010s dynamic**



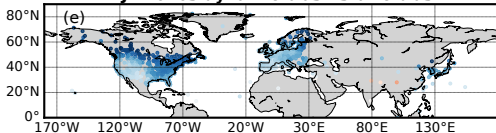
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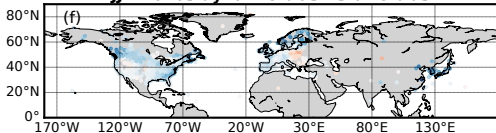
**JJA 2010s static**



**DJF 2010s dynamic bias - static bias**



**JJA 2010s dynamic bias - static bias**

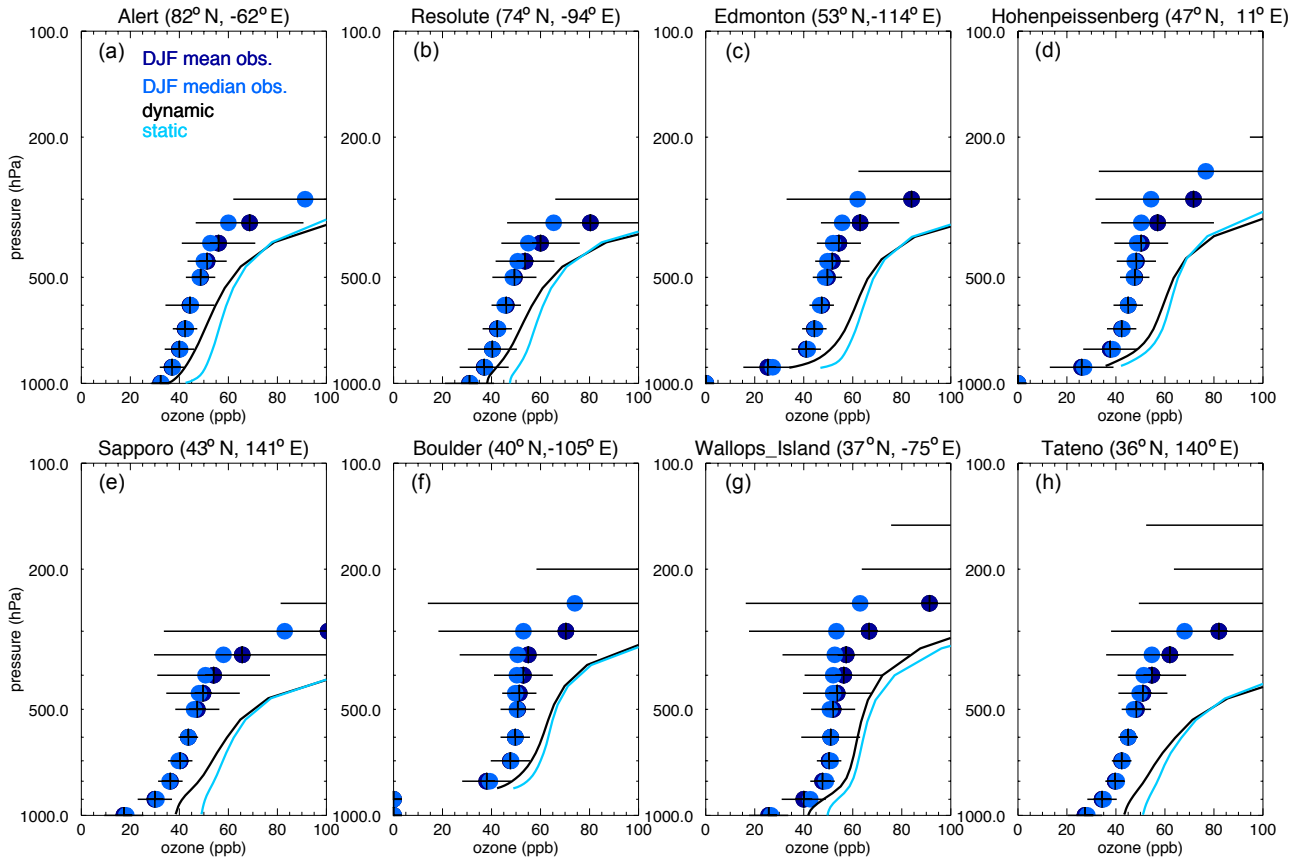


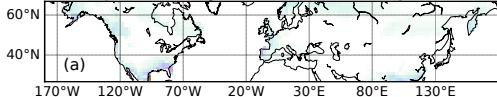
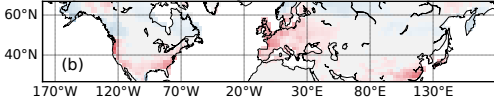
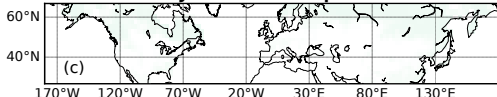
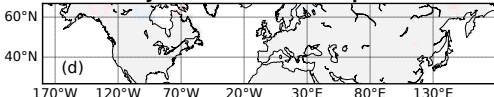
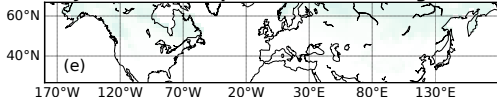
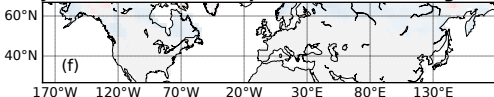
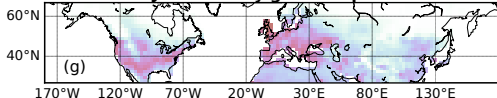
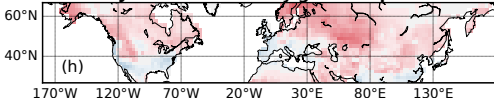
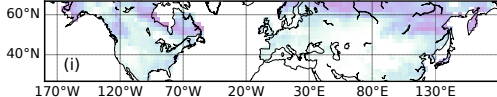
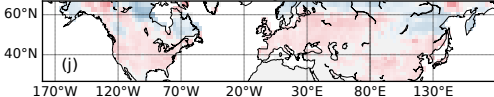
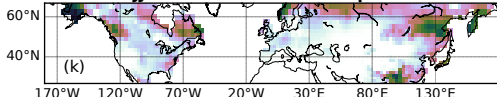
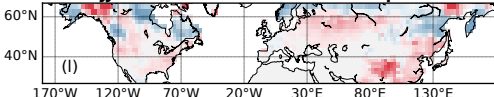
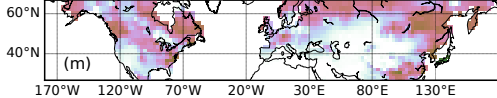
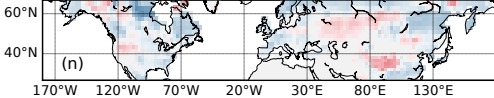
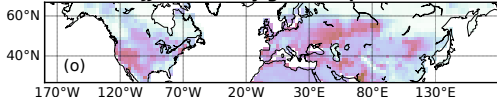
surface ozone bias (ppb)



gas difference (ppb)

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**DJF 2010s dry and wet cuticular uptake****DJF 2010s-2010s dry and wet cuticular uptake****DJF 2010s stem uptake****DJF 2010s-2010s stem uptake****DJF 2010s snow uptake to cuticles and ground****DJF 2010s-2010s snow uptake to cuticles and ground****DJF 2010s dry ground uptake****DJF 2010s-2010s dry ground uptake****JJA 2010s dry cuticular uptake****JJA 2010s-2010s dry cuticular uptake****JJA 2010s wet cuticular uptake****JJA 2010s-2010s wet cuticular uptake****JJA 2010s stomatal uptake****JJA 2010s-2010s stomatal uptake****JJA 2010s dry ground uptake****JJA 2010s-2010s dry ground uptake**